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THE EFFECT OF HEMOLYTIC STREPTOCOCCI AND THEIR PRODUCTS ON LEUCOCYTES

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Knowledge of the nature of the injury done to the tissues and wandering cells of the host by invading organisms and their products is fundamental to progress in the treatment of infectious diseases. The injury that may be done by hemolytic streptococci to the leucocytes, which are much concerned in the combat between host and invading bacteria, is the subject of the investigation here reported.

REVIEW OF LITERATURE

In the literature on infectious diseases there are many references to a decline in phagocytic activity against the specific infecting organism in fatal cases. In most of the investigations no inquiry was made to determine which of the two chief factors involved was at fault—antibody content or leucocytic efficiency; and those who have ascribed a decrease of phagocytic activity to injury of the leucocytes have generally not determined what factor in the bacterial product was responsible for the injury.

Cross found no decrease in phagocytic activity against any bacteria not concerned in the primary infection, even in the late stages of fatal disease. His results, suggesting that the nonspecific factor, the leucocytic efficiency, is not at fault in fatal cases, are at variance with the results of other investigators who have shown that streptococci and staphylococci disintegrate leucocytes.

As long ago as 1894 Van de Velde found a substance capable of disintegrating leucocytes in the exudate obtained by injecting staphylococci into the pleural cavity of rabbits. He found that this substance was destroyed at about 58° C. From this fact he concluded that it was albuminous. He gave it the name "leucocidin." He was able to demonstrate the action of leucocidin in test tube experiments as well as in vivo. Later Neisser and Wechsberg studied the staphylococcal leucocidin and concluded it was not the same as hemolysin, because the two toxic substances did not appear and disappear under the same conditions.

The following review of the literature on the production of substances harmful to leucocytes by streptococci reveals uncertainty and misunderstanding. No definite facts comparable to the facts known about staphylococcal leucocidin are established.

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M'Leod (1915) reported that, in streptococcal septicemia accompanied by marked hemolysis, the protoplasm of the leucocytes is completely disintegrated. He was unable, however, to show the action of the leucocidic substance in vitro with filtrates of streptococcus cultures. Many years later he returned to the problem, and reported (Channon and M'Leod, 1929) that, if the filtrate from a young streptococcus culture were concentrated to one-fourth of its original volume, a toxic substance capable of disintegrating leucocytes could be demonstrated in the concentrate.

Channon and M'Leod call attention to the fact that no evidence has been obtained to show that the cytolytic effects of streptococci on red

cells and leucocytes are due to different toxic substances.

Levaditi (1918) reported that he found incontestable proof that streptococci as well as staphylococci possess the power of destroying white cells in vitro. He was unable, however, to demonstrate leucocidic substance in filtrates of young or old cultures, or in extracts of dead microbes, or in macerated living streptococci. He concluded, therefore, thats treptococcal leucocidic substance is connected with the vitality of the microbes and is active only when they come in contact with leucocytes.

Nakayama (1920) believed that he could demonstrate a leucocidic substance in streptococcus culture filtrates. He used two tests for the vitality of the leucocytes: (1) The observation of ameboid movements; (2) the bioscopic tests devised by Neisser and Wechsberg to demonstrate staphylococcal leucocidin. In this test the capacity of the leucocytes for the reduction of methylene blue is taken as a measure of their vitality. Nakayama's results are confused by the use of glucose in the culture medium. It will be pointed out further on that acids are toxic for leucocytes. Hence carbohydrates, from which streptococci produce acids, should be excluded from experiments planned to demonstrate a toxic substance of the nature of that described by Van de Velde.

Using the bioscopic test of Neisser and Wechsberg for testing the vitality of cells, Dold (1930) reported that the streptococcal toxins in culture filtrates destroy not only leucocytes but also other tissue cells. Not all strains of hemolytic streptococci were found to produce the toxic substance, however, and a given strain sometimes would, and at other times would not, show evidence of its production.

Among the later writers on the subject, Wright and his collaborators (Wright, Colebrook, and Storer; Colebrook; and Hare) have reported experiments in which the phagocytic capacity of leucocytes from patients' blood was tested. They found that in septic infections the efficiency of the leucocytes is definitely subnormal when tested in normal serum, and that this efficiency appears to be reduced for all microbes indiscriminately.

No data could be found in the literature which would show whether or not the streptococcal toxin capable of producing a characteristic skin reaction (referred to hereafter in this paper as skin toxin) is toxic for leucocytes. Because many investigators have reported that there is no relationship between toxin production and virulence, it is generally inferred that the skin toxin does not affect leucocytes. That not all investigators accept that point of view, however, is illustrated by the following excerpt from Downie's recent paper: "From a histological study of the lesions it would appear that toxin acts by preventing phagocytosis so that the organisms can establish themselves and produce sufficient toxin to cause death of the animal.

* * The marked leucocytic accumulation at the site of intradermal injection in the toxin-immunized, as compared with the absence of such reaction in the coccus-immunized rabbits, is further evidence of the antiphagocytic action of toxin."

In 1922 the writer reported the sensitiveness of leucocytes to acids. Hydrochloric acid was found to be toxic in weak dilutions. Lactic acid caused more injury than hydrochloric, and acetic and butyric more than lactic, when all the acids were of the same H ion concentration. The effect of the acids on the leucocytes was cumulative. If the leucocytes were washed several times with an acid solution too weak to cause injury by a single washing, they absorbed the acid from the solution in each washing until finally enough had been absorbed to incapacitate them for phagocytosis.

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Since many pathogenic bacteria, including streptococci, are vigorous producers of acids, and since the acids have been shown to be injurious to leucocytes, they must be considered as one of the possible agents which may incapacitate the leucocytes during the progress of a disease caused by acid-producing bacteria.

All the literature on streptococci that has been reviewed may be summed up as follows:

- 1. Streptococci destroy the phagocytic capacity of leucocytes.
- 2. No data were found which would show the effect of the skin toxin on leucocytes.
- 3. Although streptococci produce acids which have been shown to be toxic for leucocytes, acids have not been considered as one of the toxic substances which may incapacitate leucocytes in vivo; and some investigators have complicated their experiments planned to show a thermolabile leucocidic substance by failing to eliminate acids from their medium.
- Whether or not streptococcal leucocidic substance is identical with hemolysin remains an open question.

EXPERIMENTAL WORK

In this study three methods were used to demonstrate the injurious effects of streptococci and their products on leucocytes: (1) The capacity of the treated leucocytes to ingest sensitized bacteria was determined by a modified Neufeld's technique; (2) disintegration of the treated leucocytes was observed in microscopic preparations; (3) the vitality of the treated leucocytes was determined by testing their capacity for the reduction of methylene blue (the bioscopic test of Neisser and Wechsberg).

THE PHAGOCYTIC TEST

A modified Neufeld's technique, the same as that used in the earlier study on the toxic effect of acids on leucocytes, was employed.

The fluid for the dilution of serum, for the dilution of test substances, and for the suspension of leucocytes in control tests was prepared by the addition of 1 part of Sorensen's phosphate buffer mixture adjusted to pH 7.0 to 9 parts of a 0.9 per cent sodium chloride solution. (It was found in the earlier study that a buffered solution was necessary for the protection of the leucocytes against chance contact with unfavorably acid solutions.)

H ion determinations were made with the use of standard buffer solutions and dye indicators.

A strain of hemolytic streptococcus originally cultivated from a case of erysipelas was used as "food" for the leucocytes and for the preparation of an immune serum for its sensitization. The serum was prepared by injecting a rabbit repeatedly with increasing doses of an antigen killed with formalin and thoroughly washed. It was preserved with 0.2 per cent tricresol. It had been kept in a refrigerator for about five months when these tests were made. It had a bacteriotropin titer of approximately 1:1280. The experiments were all carried out in triplicate, in low dilutions of the serum, as indicated in the protocols.

Two-tenths of a cubic centimeter of diluted serum and an equal quantity of a 24-hour broth culture of the streptococcus were placed together in 1 by 7 centimeter reagent tubes and incubated in a 37° C. water bath for 45 minutes. During the incubation the leucocyte suspensions were prepared.

Rabbit leucocytes were used. They were obtained by injecting into each pleural cavity about 5 cubic centimeters of sterile aleuronat 2 suspension on the day preceding the test.

All solutions in which the leucocytes were to be suspended were warmed to 37° C. The exudate was taken up in a solution of 1 per

¹ The bacteriotropins are called "stable opsonins" by some writers.

³The aleuronat suspension was made by adding 3 per cent starch and 5 per cent aleuronat to ordinary broth.

cent sodium citrate in physiological saline solution. About 50 cubic centimeters of the citrate solution was used for washing each pleural cavity. If the exudate was very bloody, the first fractions of the bloody washings were discarded and the later fractions were usually found to be sufficiently free from red blood corpuscles to be used in the test. Usually small particles of aleuronat or small clots of fibrin or blood were washed out with the exudate. They sank to the bottom of the container and were disposed of by decanting the supernatant suspension into a fresh container, mixing the leucocytes from the two pleural cavities.

A 12-cubic centimeter portion of the leucocyte suspension was placed in each of as many centrifuge tubes as there were substances to be tested. The suspensions were centrifugated for four minutes at such a speed that the majority of leucocytes were thrown to the bottom of the tube, leaving a slightly clouded supernatant fluid (the cloudiness indicating that the leucocytes had not been subjected to a compression great enough to injure them). The supernatant fluid was poured away and the sediment was emulsified in 12 cubic centimeters of buffered saline solution or test material according to the plan of the experiment. (In the protocols this is called the "second washing.") The suspension was centrifugated again in the same manner as before. This sediment was carefully emulsified in 1.5 cubic centimeters of the test or control solution and the leucocytes were then ready for the test.

Two-tenths of a cubic centimeter of leucocyte suspension was added to each tube of sensitized bacteria. The tubes were shaken to obtain a uniform suspension, and then were returned to the water bath for further incubation. During this second incubation period the racks containing the tubes were kept in vigorous motion by an electric shaking apparatus, in order to prevent the leucocytes from sinking to the bottom of the tubes. After 45 minutes' incubation, the tubes were removed from the water bath and smears were made. Before making a smear, a uniform suspension was obtained by vigorously rolling the tube between the hands. After drying, the smears were fixed with methyl alcohol. After drying again, they were stained by submerging the slides for a few minutes in a weak solution of Bordet-Gengou's toluidine blue.

Phagocytosis by the polymorphonuclear leucocytes alone was considered in this study. A characteristic picture of the phagocytosis of bacteria which have been sensitized with immune serum shows a large percentage of those leucocytes which participate in phagocytosis crowded full of bacteria. For this reason it was impossible to

³ Bordet-Gengou's toluidine blue is made by dissolving 5 grams of toluidine blue in 100 cubic centimeters of alcohol, 500 cubic centimeters of water, and 500 cubic centimeters of 5 per cent phenol, and filtering after one or two hours. One part of stain was diluted with two parts of water for staining the smears.

count the number of cocci ingested as is commonly done in the opsonic test. Twenty-five polymorphonuclear leucocytes in each smear were examined, and the presence of bacteria was recorded in terms of percentage. It was observed that those leucocytes that were agglutinated generally contained more bacteria than the isolated leucocytes. Therefore, if there had been a clumping of the leucocytes, about one-half of the number counted was chosen from one or more groups and the remainder were counted from the isolated leucocytes. Record was kept of the percentage of phagocyting leucocytes, and also of the percentage of leucocytes containing more than 10 cocci. They were tabulated in terms of leucocytes "filled" with bacteria.

Description of the toxins.—The streptococcal skin toxin used in these experiments was prepared by Surg. M. V. Veldee, of the National Institute of Health, for his own experimental work. The strain used for the preparation of the toxin was the well-known "N. Y. 5," originally cultivated by Doctor Dochez from a case of scarlet fever. The organism was grown for 89 hours at 37° C., in Douglas tryptic digest medium, as described by Watson and Wallace. The filtered toxin was of an H ion concentration of pH 7.6. It had a toxin content of approximately 60,000 skin test doses per cubic centimeter.

Diphtherial and tetanus toxins were included in some of the tests to compare their action on leucocytes with that of the streptococcal toxin. The diphtherial toxin was a sample which had been sent to the National Institute of Health by a commercial firm. It contained approximately 500 M. L. D. per cubic centimeter for guinea pigs weighing 250 grams. The National Institute of Health standard tetanus toxin was used, diluted in buffered saline solution as indicated in the respective protocols.

The effect of streptococcal skin toxin on the phagocytic capacity of leucocytes.—The phagocytic experiments were planned so that the activity of the leucocytes exposed to streptococcal skin toxin could be compared with the activity of those exposed to several inert substances, in order to demonstrate the uniformity of the phagocytic activity of healthy leucocytes. Tetanus and diphtherial toxins served as inert substances. (Many years ago Bordet showed that diphtherial toxin does not affect leucocytes.) To demonstrate the sensitiveness of leucocytes to harmful substances, parallel tests were made with solutions of acetic acid and phenol. Acetic acid was chosen because it is one of the acids produced in the fermentation of carbohydrates by streptococci. (Langwill.)

The conditions for the parallel phagocytic tests on any given date were the same except for the one variable condition of the exposure of the leucocytes to the various test or control substances. Hence the given figures in any one protocol are comparable, but they are not comparable with the figures given in other protocols, because conditions such as abnormal temperatures to which the leucocytes might be subjected during the course of preparation of the suspension, the phagocytic efficiency of the leucocytes of the different individual rabbits, and other conditions might vary from day to day. Thus the figures for the uninjured leucocytes are markedly lower in the protocols shown in Tables 2 and 4 than in those shown in Tables 1 and 3. The conclusions to be drawn from the several protocols, however, are in agreement.

It was a surprise to find that in 0.1 or 0.2 per cent solutions of phenol there was no inhibition of phagocytosis (see Table 1). It had previously been shown that phagocytosis was completely inhibited by 0.5 per cent solution. The limit of toleration for phenol was, therefore, determined. It was found that, although a 0.2 per cent solution does not affect the activity of the leucocytes, it is completely inhibited in a 0.4 per cent solution (see Table 2), and there is only slight activity in a 0.3 per cent solution (see Table 3).

The results of all the experiments agreed in showing that under the specified conditions streptococcal skin toxin has no effect on the phagocytic activity of leucocytes (see Tables 1, 2, and 3).

Recovery of leucocytes after injury due to acid.—The sensitiveness of leucocytes to acetic acid is demonstrated in Tables 2, 3, and 4. results showing inhibition of phagocytosis by acetic acid agreed with those reported in the earlier publication.

Table 1.—Protocol of experiment showing that, under the conditions of the phagocylic test, streptococcal skin toxin, tetanus toxin, and 0.1 per cent phenol do not injure the leucocytes. (Second washing of leucocytes was in buffered saline solution; final suspension was a solution of the test or control material)

No. of the	Control or test material	pH of the so- lutions of the	1	rum t	n of imm used for a n of strep	Streptococci not sensitized but suspended in control solu- tions		
test		control or test mate- rial		1:20	1:40	1:80	Buf- fered saline solution	Normal serum diluted 1:20
1	Buffered saline solution Streptococcal toxin	7.0	-	1 68 1 48 76	56 28 64	76 36 60	12 0 16	4
3 4	Tetanus toxin 3	6.4	-	3 48 76 56 56 40 64 56	44 64 56 76 40	76 36 60 40 72 64 48 20	0 8 0 12	0

The upper figure refers to the percentage of phagocyting leucocytes.
 The lower figure refers to the percentage of leucocytes containing 10 or more coccl.
 The tetanus toxin was diluted to contain approximately 4,800 M. L. D. per cubic centimeter for 350gram guinea pigs.

TABLE 2.—Protocol of experiment showing that, under the conditions of the phagocytic test, streptococcal skin toxin, tetanus toxin, and diphtherial toxin do not, while acetic acid and 0.4 per cent phenol do, injure the leucocytes. (Second washing and final suspension of the leucocytes were in solutions of the test or control material)

No. of the test Control	Control or test material	pH of the so- lutions of the	rum	n of imm used for n of strep	but sus	Streptococci not sensitized but suspended in control solu- tions	
	Control of cess material	control or test mate- rial	1:20	1:40	1:80	Buf- fered saline solution	Normal serum diluted 1:20
	Buffered saline solution	7.0	1 44 2 20 44 16	48 16 36 16	40 20 36 16 32	10 4 13	8
	Tetanus toxin Diphtherial toxin	6.4 7.6	32 20 48 16	52 16 40 12 28	20 32 12	16 4 8	10
	Acetic acid	7.0	16 0 64 28 12	28 4 56 28	16 12 40 28	16	8
	0.4 per cent phenol	7.0	12	8	4	0	

¹⁹ See Table 1 for significance of the figures.

The tetanus toxin was diluted to contain approximately 3,000 M. L. D. per cubic centimeter for 350-

Table 3.—Protocol of experiment showing that, under the conditions of the phagocytic test, streptococcal skin toxin and diphtherial toxin do not, while acetic acid and 0.3 per cent phenol do, injure the leucocytes. (Second washing and final suspension of the leucocytes were in solutions of the control or test material)

No.	Control or test material	pH of the so- lutions of the	rum t	n of imm used for n of strep	Streptococci not sensitized but suspended in control solu- tions		
test	Control of sess material	or test mate- rial	1:20	1:40	1:80	Buf- fered saline solution	Normal serum diluted 1:20
1	Buffered saline solution	7. 0	{ 1 00 40	68	40 28	16	0
2	Streptococcal toxin.	7. 6	64 28	48 36	48 32	12	8
3	Diphtherial toxin	6.4	64 28 64 32 28 8 20	44 48 36 52 24 24 12	40 28 48 32 48 12 32	4 0	12
4	Acetie acid.	4.8	28	24 12	32	8	8
5	0.3 per cent phenol	7.0	20	24	32	Ō	4

¹ See Table 1 for the significance of the figures.

A few experiments were carried out to determine whether leucocytes readily recover from the injury caused by acetic acid. The protocol of a typical experiment is given in Table 4. The technique used for the phagocytic experiments previously described in this paper was modified for these tests. Leucocytes slightly injured so that phagocytic

³ The tetanus toxin was diluted to contain approximately 3,000 M. L. D. per cubic centimeter for 350 gram guinea pigs.

activity was only partially destroyed and leucocytes which had been exposed to amounts of acid very slightly exceeding their limit of toleration were used. Under the conditions of these experiments leucocytes washed in acetic acid of an H ion concentration of pH 4.8 showed partial destruction of phagocytic activity, and those washed in acetic acid of an H ion concentration of pH 4.6 showed almost or quite complete destruction of phagocytic activity. These and slightly greater concentrations of acid, were used in the experiments.

The leucocyte suspension in sodium citrate solution as obtained from the pleural cavities of a rabbit was divided into eight portions and centrifugated, and the supernatant fluid was poured away. The tubes were then divided into two series, A and B, of four tubes each. The leucocytes in the tubes of the series designated A were tested for phagocytic activity immediately after washing in the various control and test solutions, and those in the corresponding tubes of the B series which had been exposed to the same solutions as those of the A series were suspended in several cubic centimeters of fresh serum from a normal rabbit and incubated in a water bath at 37° C. for an hour or two and then tested for phagocytic activity. The treatment of the leucocytes in the various tubes after the first centrifugation and disposal of the sodium citrate solution was as follows:

Series A. Tube 1 (control)—(a) The leucocytes were washed in 12 cubic centimeters of buffered saline solution and centrifugated. (b) They were resuspended in 1.5 cubic centimeters of the buffered saline solution and added to the sensitized bacteria to test for phagocytic capacity.

Tube 2.—The leucocytes were treated like those in Tube 1 except that the washing and final suspension was in acetic acid of pH 4.8.

Tube 3.—The leucocytes were treated like those in Tube 1 except that the washing and final suspension was in acetic acid of pH 4.6.

Tube 4.—The leucocytes were treated like those in Tube 1 except that the washing and final suspension was in acetic acid of pH 4.4.

Series B. The four tubes of Series B were treated like the corresponding tubes of Series A through (a) and (b). (c) The suspensions were centrifugated and the leucocytes were resuspended in a few cubic centimeters of fresh rabbit serum and incubated for an hour or two in a water bath at 37° C. (d) The suspensions were centrifugated again and the supernatant serum was poured away. (e) The leucocytes were resuspended in 1.5 cubic centimeters of buffered saline solution, and this suspension was added to sensitized bacteria to test the phagocytic capacity of the leucocytes. All the tests were carried out in triplicate, with the same strain of streptococcus sensitized with the same increasing dilutions, all low, of the same high titered homologous serum used in the experiments recorded in Tables 1, 2, and 3.

Table 4 shows that leucocytes whose phagocytic capacity had been slightly injured by washing in acetic acid of pH 4.8 were restored to their usual activity (as compared with leucocytes washed in buffered saline solution) by incubation in fresh serum. Leucocytes whose phagocytic activity had been almost completely inhibited by washing in acetic acid of pH 4.6 were also restored to their usual activity by incubation in fresh serum. On the other hand, the leucocytes which had been washed in acetic acid of pH 4.4 were so badly injured that incubation in fresh serum had no effect on them. The experiment was repeated several times with similar results.

Table 4.—Protocol of experiment showing the effect of incubation in normal serum on leucocytes which have been injured by acetic acid

[The triplicate sets of figures show leucocytic activity for bacteria sensitized with three different low dilutions of homologous immune serum]

No. of the tube containing the leucocyte suspension	Trestment of the leucocytes	(Activity was to ately	after wa	medi-	Series B (Activity of leucocytes was tested after wash- ing in control or test solutions and then incu bating for an hour infresh serum)		
	{(Control) washed in buffered saline solution pH 7.0. Washed in acetic acid, pH 4.8	{ 132 12 16 12 12 12 4 4 4	40 20 20 12 4 0	36 20 16 4 4 0 4	40 20 28 20 32 20 4 0	32 16 28 16 20 16 0	28 16 44 20 32 20 4

¹¹ See Table 1 for the significance of the figures.

It would be impossible to duplicate in a test tube experiment the injury done to leucocytes by the acids produced by the bacteria in a focus of infection. The body fluids are sufficiently buffered so that the circulating blood never reaches an H ion concentration low enough to affect the leucocytes. But due to their strong affinity for acids it appears possible that the leucocytes accumulated at the site of infection may gradually take up the acids until their limit of toleration is reached and phagocytic capacity is finally crippled. There would be a continuous absorption of dilute acids, and at the same time there would be a more or less continuous restoration to a healthy condition, dependent on the flow of blood through the focus of infection. The results of the experiments suggest that leucocytes slightly injured by acid may be restored to their usual activity if there is a good circulation of blood in the focus of infection; whereas if the blood supply is deficient, the leucocytes may become injured beyond recovery.

THE DISINTEGRATION OF LEUCOCYTES BY STREPTOCOCCI 4

An attempt was made to demonstrate the disintegration of washed leucocytes in the presence of washed streptococci by changes in the H ion concentration of the saline solution in which they were suspended. A slight increase of H ions was sometimes indicated but this method of detecting the disintegration of leucocytes was abandoned because there were too many complicating factors, chief among which was the increase of H ions, due to the autolysis of the leucocytes.

When washed leucocytes and washed streptococci were suspended together in physiological saline solution, the disintegration of the leucocytes could be observed in microscopic preparations. technique described for the phagocytic test was used for obtaining and washing the leucocytes and for the preparation of slides for microscopic examination. It was necessary to wash the streptococci rapidly because their capacity for attacking the leucocytes was quickly injured by saline solution. The culture was centrifugated, the supernatant fluid was removed, and a few cubic centimeters of saline solution were allowed to flow gently over the sediment without disturbing The wash fluid was removed and the sediment was emulsified in a small quantity of saline solution, making a heavy suspension, which was immediately added to a suspension of washed leucocytes. Under these conditions there was practically no phagocytosis. Smears prepared after 2, 3, or 4 hours' incubation showed definite disintegration of leucocytes, as compared with control suspensions without streptococci, or with streptococci killed by heat. The disintegrated leucocytes appeared as faintly stained forms, without demonstrable nuclei. After longer incubation the leucocytes in the control tubes underwent similar changes, due to autolysis.

The lysis of leucocytes by living streptococci could be more readily demonstrated if broth instead of saline solution were used for washing the streptococci and leucocytes and for the final suspension in the experiment just outlined. Leucocytes suspended in broth do not autolyze for many hours. Hence there was a definite contrast between the disintegrated leucocytes in the suspensions with living streptococci and the healthy leucocytes in the control suspensions. The contrast was marked after three or four hours' incubation. After 21 hours' incubation no recognizable leucocytes could be found in smears of the growing streptococcus cultures, whereas those in smears from the control tubes were fairly well stained.

If leucocytes were suspended in a filtrate of broth culture of hemolytic streptococci they retained their staining properties as well as if suspended in broth. Hence it may be stated that no demonstrable toxic substance is excreted into broth by growing streptococci

⁴ The scarlet fever strain known as Dick I was used in these tests.

when judged by the effect of the filtrate on the staining properties of the leucocytes.

THE BIOSCOPIC TEST

The bioscopic test of Neisser and Wechsberg is the most delicate test for determining whether leucocytes have been injured. In this test the vitality of the leucocytes is measured by their capacity for reducing methylene blue to the colorless reductant.

The test was carried out as follows: A suspension of washed leucocytes was obtained in the same manner as that employed for the phagocytic test. One-half of a cubic centimeter of heavy leucocyte suspension (the yield from one rabbit in 6 or 8 cubic centimeters of broth) were added to 2 cubic centimeters of the test or control solution in 1 by 7 millimeter reagent tubes, with 2 drops of 0.5 per cent aqueous solution of methylene blue. A uniform suspension was obtained by drawing the mixture into a pipette; then the contents of the tubes were covered with a layer of 0.5 cubic centimeter of liquid petrolatum. Control tubes were always set up without leucocytes to show that there was nothing in the test fluid which would bring about the reduction of methylene blue. The rack of tubes was placed in a 37° C. water bath, and readings were made at intervals up to two hours.

The demonstration of a toxic substance by the bioscopic test.—The bioscopic test was used to demonstrate substances injurious to leucocytes in scarlet fever skin toxin, in filtrates of young broth cultures of hemolytic streptococci, and in filtrates of cultures in broth with various additions. Kidney tissue, blood serum, washed leucocytes or washed erythrocytes from rabbits were added to broth at different times to determine their possible influence on the production of leucocidic substances. The tests were usually made with filtrates of 24-hour cultures, although it was found that the results were the same when tests were made with filtrates of older cultures. The strain known as Dick I was used in the preparation of filtrates for some of the tests and a strain, No. 663, freshly isolated from the throat in a case of scarlet fever was used for the preparation of filtrates for other tests. The two strains gave the same results.

Table 5.—Protocol of bioscopic tests showing that a trace of toxic substance is excreted into broth by growing hemolytic streptococci

	Incubated for—							
Leucocytes were suspended in—	30 minutes	1 hour	2 hours					
Broth, pH 7.0. Bearlet-fever toxin Filtrate of broth culture.	Complete reduction	Complete reduction.						
Broth with acetic acid, pH 4.8	No reduction	No reduction	No reduction					

Table 6.—Protocol of bioscopic tests showing that the addition of kidney tissue, blood serum, or washed leucocytes does not influence the production of toxic substance, whereas it is produced abundantly in broth containing washed erythrocytes.

A CONTRACTOR OF THE PARTY OF TH	Incubated for—						
Leucocytes were suspended in—	30 minutes	1 hour	2 hours				
Broth Filtrate of broth culture Filtrate of culture in broth plus kidney tissue. Filtrate of culture in broth plus blood serum. Filtrate of culture in broth plus washed leucocytes. Filtrate of culture in broth plus washed erythrocytes.	Complete reduction	Complete reductiondodododo	No reduction.				

The results of repeated tests are summarized in Table 5. Broth adjusted to a reaction of pH 4.8, by the addition of acetic acid, was included among the test substances to compare the effect of the toxic substance in question with that of a known toxic substance. acetic acid completely inhibited reduction. The table shows that the leucocytes suspended in broth completely reduced the methylene blue in 30 minutes, whereas leucocytes suspended in scarlet fever skin toxin or in filtrates of young broth cultures partially reduced the methylene blue in 30 minutes, with complete reduction within an hour. This delay of reduction always occurred in tests with leucocytes suspended in the toxin or filtrates of young broth cultures, as compared with leucocytes suspended in broth. The data thus obtained with the bioscopic test show that there is a trace of leucocidic substance produced in broth by growing hemolytic streptococci. It may be recalled that no trace of this leucocidic substance could be detected by the phagocytic test, nor in microscopic preparations of treated leucocytes.

An effort was made to find some substance available to strepto-cocci when they grow as parasites which might promote an excretion into the medium of the leucocidic substance. The data are presented in Table 6, which shows that the addition to broth of kidney tissue, blood serum, or washed leucocytes did not influence the production of leucocidic substance in the culture medium. In repeated tests the delay of reduction was the same for leucocytes suspended in filtrates of cultures grown in these media as for leucocytes suspended in filtrates of broth culture. On the other hand, the addition of washed erythrocytes to the broth markedly promoted the production of a toxic substance. There was no reduction of methylene blue during the two hours of observation when leucocytes were suspended in filtrate of culture in broth to which washed red cells (10 per cent of red cell suspension in which the washed cells were supsended in broth to make the original volume of blood) had been added.

⁵ This was the same sample of toxin which was used in the phagocytic tests. It had a titer of approximately 60,000 skin-test doses per cubic centimeter.

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The enhanced production of leucocidic substance in broth plus washed red cells was confirmed in microscopic preparations of leucocytes suspended in the filtrate of such a culture, as compared with those suspended in broth or filtrate of broth culture. There was definitely a more rapid disintegration of leucocytes suspended in the filtrate of culture in broth plus washed red cells than in the control tubes. Thus it was demonstrated by the observation of disintegration of leucocytes in microscopic preparations as well as by the bioscopic test, that a leucocidic substance is produced by streptococci from red blood cells.

Does serum contain an agent for the neutralization of the leucocidic substance? Bioscopic tests were carried out to determine whether normal or immune serum contains an agent to neutralize the leucocidic substance. In these tests 0.5 cubic centimeter of serum was added to 1.5 cubic centimeters of the test or control fluid, and the mixture was incubated for an hour and a half; then a suspension of leucocytes and methylene blue were added as for the bioscopic tests previously described. These tests were carried out with the recently isolated scarlet fever strain of streptococcus used in previous experiments (No. 663), and with homologous immune serum of high agglutinating titer prepared with formalin killed antigen (2 serums) or with one dose of living antigen following a course of treatment with killed antigen (1 serum).

Normal serum is a better medium than broth to maintain the vitality of leucocytes, as can be demonstrated by the slightly more prompt reduction of methylene blue in broth plus 25 per cent of serum than in broth. Neither the normal nor the immune serum could be shown to enhance the reduction of methylene blue by leucocytes exposed to the leucocidic substance further than the slight advantage which was given by adding serum to the broth control. There was, therefore, no demonstrable specific neutralizing agent in either the normal or the immune serum.

THE NATURE OF THE LEUCOCIDIC SUBSTANCE

The thermolability of the substance toxic for leucocytes in the filtrate of culture in broth plus red cells was determined by means of the bioscopic test. The thermolability of the trace of leucocidic substance in scarlet fever toxin and in filtrates of young broth cultures was also determined by the same method and was found to be identical with that of the stronger leucocidic substance produced at the expense of erythrocytes. Presumably the same leucocidic substance is produced under the varying conditions. Temperatures of 37° C. for a day or under 56° C. for one hour do not affect it. There is slight destruction at 56° for one hour, and more with increasing temperatures up to 75° for one hour, at which temperature destruction

is almost complete. No trace of the leucocidic substance could be found after heating at 85° C. for one hour.

The thermolability of the leucocidic substance produced by hemolytic streptococci as reported here agrees with Van de Velde's leucocidin. He stated that the staphylococcal leucocidin is destroyed at about 58° C. There is, however, an objection to the application of the term "leucocidin" to the leucocidic substance produced by streptococci. Several authors (Eijkman; M'Leod and Govenlock; Rogers) have reported that streptococci as well as other bacteria produce a thermolabile substance which inhibits the growth of the homologous organism or other bacteria. This substance has been called "bactericidin." There is no evidence at hand to show whether or not the so-called bactericidin is identical with the leucocidic substance.

The injury done to leucocytes by the thermolabile toxic substance is quite different from that done by acid. Leucocytes injured beyond recovery by acid retain their morphology and staining properties, whereas leucocytes injured by the thermolabile toxic substance are disintegrated.

IS THE LEUCOCIDIC SUBSTANCE IDENTICAL WITH HEMOLYSIN?

Two lines of evidence are offered to show that the leucocidic substance produced by streptococci is not identical with hemolysin:
(1) They differ in thermolability; (2) under certain conditions of growth the production of the leucocidic substance is enhanced, whereas under those same conditions hemolysin production is inhibited.

According to M'Leod and M'Nee hemolysin is destroyed by heating a few hours at 37° C. Their observations on the extreme theremolability of the streptococcal hemolysin were confirmed in this study. Hemolysin was destroyed by heating overnight at 37° C. or by heating one hour at 45° C.

Since the leucocidic substance is uninjured at 37° C. for a day, or at 45° for one hour, a filtrate of broth culture containing a vigorous hemolysin can be heated to destroy all the hemolysin without injuring the trace of leucocidic substance which it contains.

If the leucocidic substance were identical with hemolysin there should be an evident correlation between the vigor of action on the two types of blood cells manifest by filtrates of cultures grown under various conditions. A filtrate containing strong hemolysin should also contain strong leucocidic substance and vice versa. Hence if leucocidin and hemolysin were identical, there should be a much stronger content of hemolysin in the filtrate of culture in broth plus red cells than in the filtrate of broth culture, for it was shown (Table 6) that the filtrate of culture in broth plus red cells contains definitely

more leucocidic substance than the filtrate of broth culture. The facts, however, are contrary to that supposition. Experiments were carried out which showed that erythrocytes added to broth not only fail to enhance the production of hemolysin by Streptococcus scarlatinae (the "Dick I" strain was used), but they even inhibit its production as compared with the production of hemolysin in broth without red cells.

Table 7.—Protocol showing that erythrocytes in broth culture of Streptococcus scarlatinae interfere with the production of hemolysin

Tube No.			Hemolysis as de- termined by ap- pearance of tubes before centrifu- gation	Erythrocytes re-	Color readings after the cryth- rocytes were he- molysed in 10 cubic centimeters of water
1	0.4 cubic centi- meter.		No hemolysis	0.2 cubic centi- meter sediment.	Deep red.
2	0.1 cubic centi- meter.	Broth (control)	do	0.05 cubic centi- meter sediment.	Pale red.
3	0.4 cubic centi- meter.	Filtrate of broth	Complete hemo-	Slight colorless sediment.	Very faint tinge of color.
4	0.1 cubic centi- meter.	culture.	do	No sediment	No color.
8	0.4 cubic centi- meter.	Filtrate of culture in broth+eryth-	Readings could not be made.	0.16 cubic centi- meter sediment.	Color is almost as deep as in (1). The distinction is questionable.
. 6	0.1 cubic centi- meter.	rocytes.		0.025 cubic centi- meter sediment.	Color is not quite so deep as in (2).

The usual color test for hemolysin was not applicable to its determination in filtrates of cultures in broth containing erythrocytes, because the red color of the filtrate made comparative readings impossible in the final test. Hence the amount of hemolysis in the various experimental fluids was determined by measuring the amount of erythrocytes remaining. In the first experiment to compare the hemolysin content of a filtrate of streptococcus culture in broth with that in broth to which erythrocytes (rabbit) were added, the test for hemolysin was made with both rabbit and human erythrocytes, with identical results. There was very little, if any, hemolysis in the filtrate of culture in broth plus erythrocytes, whereas vigorous hemolysis occurred in the filtrate of broth culture. The experiment was repeated and the results are given in Table 7. Cultures were grown overnight in broth, and in broth containing the washed erythrocytes from 10 cubic centimeters of rabbit blood in 50 cubic centimeters of broth. After filtration, 10 cubic centimeters of the various test fluids were measured into graduated centrifuge tubes, and washed rabbit erythrocytes were added. Control tests were made in broth. To one series of tubes 0.4 cubic centimeter, and to another series 0.1 cubic centimeter of suspension of washed erythrocytes was added. tubes were incubated for four hours in a 37° C. water bath, then were transferred to the refrigerator. On the following day, color readings

were made on tubes for which that was possible, then the tubes were centrifugated and the amount of sediment in the tubes was recorded. The supernatant fluid was removed and 10 cubic centimeters of water were added to each tube. After complete hemolysis had occurred, color readings were made again.

The data recorded in Table 7 show that there was only a minute quantity of hemolysin in the filtrate of culture containing erythrocytes, whereas there was abundant hemolysin in the filtrate of broth culture. A comparison of Tables 6 and 7 leads to the conclusion that the leucocidic substance is not identical with hemolysin, because the addition of erythrocytes to broth culture promotes the production of the leucocidic substance, whereas it inhibits the production of hemolysin.

IS THE LEUCOCIDIC SUBSTANCE IDENTICAL WITH SKIN TOXIN?

The thermolability of the leucocidic substance is about the same as that of the scarlet fever skin toxin. Hence, thermolability determinations gave no information as to the unity or duality of the toxic material. Evidence that the leucocidic substance is not the skin toxin was obtained, however, from the irregularity of the ratio of the two substances in various filtrates. It was noted in previous experiments that the delay in reduction of methylene blue by leucocytes was always the same, giving evidence of only a trace of leucocidic substance whether the test was made with skin toxin of a titer of 60,000 skin test doses or with filtrates of 24-hour cultures of any one of the three strains of streptococci used in the tests. Yet the "N. Y. 5" strain is known to produce two or three times as much skin toxin as the "Dick I" strain.

Table 8.—Protocol of bioscopic tests showing that concentrated skin toxin contains less leucocidic substance than the unconcentrated toxin

	Reduction after incubation for —									
Leucocytes suspended in—	20 minutes	30 minutes	35 minutes	40 minutes	45 minutes					
Broth	Complete Nonedo	Considerabledodo	Almost complete.	Completedo						
s. t. d. Unpurified toxin, 60,000 s. t. d. Filtrate of culture No. 663 Filtrate of "Dick I" culture	do	Slightdodo	Considerabledodo	Almost complete.	Do.					

A purified and concentrated preparation of skin toxin offered material for more decisive comparative tests. This toxin, prepared with the "N. Y. 5" strain, was purified and concentrated by precipitations with acetone, alcohol, acetic acid, and alcohol, respectively. The final product contained approximately 150,000 skin test doses

per cubic centimeter. Its H ion concentration was pH 8.2. A portion of the sample was adjusted to pH 7.6 by the addition of a trace of dilute acetic acid, and tests for leucocidic substance were made with both portions. Parallel tests were made with filtrates of 24hour broth cultures and with the unpurified unconcentrated skin toxin used in previous experiments. A protocol of one of the experiments is given in Table 8. After the beginning of reduction of the methylene blue, readings were made every five minutes in order to detect even slight differences in the amount of leucocidic substance present in the fluids under observation. The two samples of purified toxin of slightly different H ion concentration behaved exactly alike, and the unpurified toxin behaved exactly like the two filtrates of young streptococcus cultures. Reduction was more prompt in the two samples of purified toxin than in the sample of unpurified toxin, although the purified toxin contained two and onehalf times as many skin-test doses of toxin per cubic centimeter as the unpurified sample. The experiment was repeated with similar The results of these experiments indicate that the leucocidic substance is not identical with skin toxin.

SUMMARY

The results of the experiments may be summed up as follows:

1. Leucocytes are injured by acid. If the injury is not too great, they may be restored to a healthy condition by bathing in blood serum.

2. In filtrates of broth cultures of Streptococcus scarlatinae there is a trace of a substance toxic for leucocytes which can be detected by the bioscopic test, but not by the phagocytic test nor by the deterioration of cells as shown in stained microscopic preparations.

3. The addition of kidney tissue, blood serum, or washed leucocytes to broth cultures does not increase the production of the leucocidic substance. On the other hand, the addition of washed erythrocytes to broth cultures definitely promotes its increase.

4. The thermolability of the trace of leucocidic substance in filtrate of broth culture is the same as that of the more abundant leucocidic substance in filtrate of culture in broth plus erythrocytes. Presumably the two substances are identical.

5. A specific neutralizing agent for the leucocidic substance could not be demonstrated in normal or immune serum.

6. Two lines of evidence are offered which show that the leucocidic substance is not identical with hemolysin.

(a) They differ in thermolability.

(b) There is no correlation of toxicity for the two types of blood cells manifest by filtrates of cultures grown under varying conditions.

The decrease of leucocidic substance in purified and concentrated skin toxin indicates that leucocidic substance and skin toxin are not identical.

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RAT-FLEA SURVEY OF THE PORT OF ST. THOMAS, VIRGIN ISLANDS

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Available sanitary records of the Virgin Islands do not show that epidemics of plague have ever occurred in any of this group of the West Indies. In view of the fact that epidemics of plague have occurred in neighboring islands, a rat-flea survey of the principal port of the Virgin Islands, St. Thomas, was undertaken to determine the infectibility of this port with plague, as indicated by the *cheopis* index.

During the 1921 epidemic of plague in Porto Rico a strict quarantine was maintained in the Virgin Islands against all Porto Rican ports. The nearest Porto Rican port is only 40 miles from St. Thomas. Fortunately, shipping at that time between these islands consisted mainly of sailing vessels which usually did not dock at St. Thomas, but lay at anchor in the harbor. Nevertheless, quarantine regulations to prevent the introduction of plague were strictly and successfully enforced.

METHOD OF SURVEY

The procedure of the survey was based on similar methods used in New York (1), San Juan (2), and Norfolk (3). Rats were captured alive in cage traps and brought to the quarantine office with the cage uncovered, being handled gently to guard against dislodging any fleas from the rats.

The rats were then killed with chloroform and the fleas collected in accordance with the ingenious method devised by Hasseltine (3) in the Norfolk survey of 1927-28. A small box with a hinged glass top was used for chloroforming the rats. In one end of the box a round hole was cut, which could be closed by a sliding partition. The box was lined with white paper. The cage trap containing the live rat was placed so that the hole for the rat's egress from the trap coincided with the hole in the end of the box. The rat, attempting to escape from the trap, usually went into the box of his own volition. The partition was then slid over the hole, the hinged glass top slightly raised, and gauze saturated with chloroform introduced. The rat when dead, as observed through the window, was removed and combed for fleas. The box was also shaken out to obtain any fleas that might have become dislodged from the rat. No rats escaped.

The fleas were preserved in 95 per cent alcohol and sent to the New York quarantine station for identification of species.

The survey began July 1, 1929, and ended June 30, 1930, being carried on entirely by the regular personnel of the U. S. quarantine station at St. Thomas. During the first four months of the survey the

daily average number of traps was 28; during the last eight months the daily average number was 51.

The town was divided into four zones for purposes of trapping. Zone 1 consisted entirely of the docks where the large vessels are berthed. The dock area is about three-quarters of a mile from the town proper, lying on the opposite side of the harbor, and connected overland by a road skirting the harbor and traversing marshy open land. Zone 2 consisted of all the water front of the town proper. Here the water is shallow, and only sloops and similar small vessels can tie up to the short docks of wood or concrete. The business district skirts this water front. Zone 3 also lies on the water front, but at the extreme western end of the harbor, and comprises a small fishing village, lying about one-half mile from the town itself. Zone 4 consists of the residential district and is made up of three hills sloping upwards rather sharply from the low lying water front and business district.

The docks of zone 1, where the large ships are tied up, are of concrete. The warehouses are constructed of concrete and metal, with a concrete floor and foundation. They afford practically no rat harborage. The buildings of zone 2, are of all types of construction and afford ample rat harborage, as do those of zone 3. The buildings of the residential district are made up of some dwellings built largely of stone, concrete, and masonry, interspersed with others which range from 2-story frame dwellings to mere shacks.

In the vicinity of St. Thomas the soil is hard and rocky, with scant vegetation.

DISTRIBUTION OF RATS

The total number of trap-days was 15,755; the daily average number of traps was 43. During the 365 days of trapping, 312 rats were caught, and a total of 2,113 fleas retrieved. Of the 312 rats, 309 were identified as Rattus alexandrinus, and 3 as Rattus rattus. None of the species Rattus norvegicus was found.

The greatest number of rats were taken in zones 2 and 4, where harborage was found to be most ample. Only three rats were captured in zone 1, which comprised the area of concrete docks and ratproof warehouses.

Entire absence of the species Rattus norvegicus seemed unusual; but this is probably due to two factors. One of these is the absence of suitable harborage for this species. The soil is extremely hard and rocky, precluding much possibility of burrowing refuges. The sewers, most of which are open, are of concrete and masonry, running for comparatively short distances downhill to the sea. The second, and probably the most important factor, is the presence of the mongoose, which overruns the island and is the rat's natural enemy. The

presence of the mongoose and the lack of suitable harborage have probably caused the elimination of all of the rat species not adapted to life in trees or houses.

TABLE 1 .- Distribution of rats and fleas by months

		Rat	ttus			Rats	2.5		Fleas			
	Total rats	alexa	ndri- us		tus	per hun- dred traps days	X. cł	neopis		canis felis		Che- opis index
		Male	Fe- male	Male	Fe- male	per month	Male	Fe- male	Male	Fe- male	Total	
1929												
July	18	14	3	1		2.0+	95	66		*****	161	8.90
August	16	6	10			1.9-	95	68			163	10. 10
September	17	10	7			2.0+	93 63	61			154	9. 0
October	18	13	5			2.0+	63	50			113	6. 27
November	27	18	9			1.7+	46	33	2	1	82	2.92
December	19	8	11			1.2+	45	. 51			96	5.00
January	22	10	12			1.4+	49	34			83	3. 60
February	23 31	ii	13 20			21+	92	124	1		217	7.0
March	32	19	12	1		2.0+	105	160			265	8, 28
April	36	16	20			2.3+	95	133		1	229	6. 33
May	39	20	18		1	2.4+	119	153			272	6, 99
June	36	18	18			2.3+	130	148			278	7. 70
Total	312	163	146	2	1	• 1.9	1,027	1, 081	8	2	2, 113	• 6.75

[·] Average.

TABLE 2 .- Distribution of rats and fleas by zones

Zone				Total				
	Total number of rats caught	X. eh	eopis	Ct. cani	s or felis	Total	Total number of fleas per rat	number of X. che opis per
	Caught	Male	Female	Male	Female	Total	por rai	rat
1 2 3	3 134 42 133	8 462 96 495	9 485 118 435	1 2	1	17 949 214 933	5. 7 7. 00 5. 00 7. 00	5. 7 7. 09 5. 00 7. 00
Total	312	1,061	1,047	3	2	2, 113	6.77	6.7

Table 1 shows the distribution of rats and fleas by months, Table 2 presents the distribution by zones, and Chart 1 shows the relations of temperature, rainfall, "rat take" by months, and *cheopis* index. As no data of relative humidity were obtainable, the amount of rainfall by months was substituted for this factor.

DISTRIBUTION OF FLEAS

A total number of 2,113 fleas was recovered from 312 rats. Of this number 2,108, or 99.7 per cent, were identified as Xenopsylla cheopis, and 5, or 0.3 per cent as Ctenocephalus felis or canis. Relative proportions of male and female are shown in Table 1. The average number of fleas per rat was 6.7, and as Xenopsylla cheopis constituted

99.7 per cent of the fleas, the *cheopis* index (4) for all practical purposes may be taken as the same figure, 6.7.

The St. Thomas cheopis index of 6.7 is only slightly below the cheopis index of 7.05 of San Juan (2), where plague has occurred within the past 9 years. The index is higher than that of New Orleans and other ports of the continental United States.

Factors that influence the prevention of the introduction of plague into this port are the practically rat-proof docks and warehouses, the distance of these docks from the main body of the town, and the character of the shipping entering the port. St. Thomas is largely a bunkering port, the majority of vessels being in port for a few hours only to obtain bunker coal or fuel oil. The greater number of vessels arriving from ports plague-infected, or recently plague-infected, are

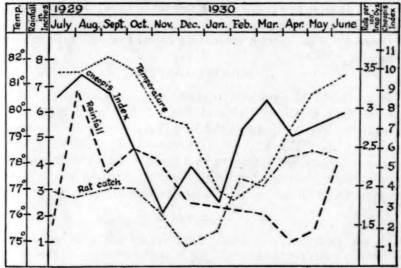


CHART 1.—Graphs showing temperature, rainfall, number of rats caught per 100 trap days, and cheopis index

laden with such cargoes as nitrates, ores, etc., which do not attract rats. Most of these vessels enter under provisional pratique and are required to breast off 4 feet from the dock, apply standard rat guards on all lines, and raise gangways at night. As soon as they have finished coaling, they depart. In the case of vessels from ports badly infected with plague, in addition to these precautions such vessels are allowed alongside the dock only during daylight hours and are kept under strict surveillance.

SUMMARY

1. A rat-flea survey of the port of St. Thomas, Virgin Islands, from July 1, 1929, to June 30, 1930, resulted in the capture of 312 rats, from which 2,113 fleas were taken.

2. Of the 2,113 fleas, 2,108, or 99.7 per cent, were identified as Xenopsylla cheopis, and 5, or 0.3 per cent, as Ctenocephalus canis or felis.

3. On the basis of the figures obtained, the average rat-flea index for the period was 6.7, which was approximately the Xenopsylla

cheopis index.

4. The cheopis index was high throughout the year, but relatively highest during the summer months (March to September, inclusive) and varied in direct relation to temperature and rainfall.

5. Rattus alexandrinus was found to be the predominating rat.

None of the species, Rattus norvegicus, was found.

6. It would seem that, should plague be introduced, it would spread

rapidly, as all conditions appear favorable for its propagation.

7. All possible precautions are being taken to prevent the introduction of plague by shipping, and the local sanitary authorities, advised of the result of the survey, are making efforts toward a rat-eradication campaign.

ACKNOWLEDGMENTS

It is desired to express appreciation and acknowledge indebted ness to Medical Director Carroll Fox, of the United States Public Health Service, in charge of the New York quarantine station, and to Surg. C. L. Williams, of the United States Public Health Service, in charge of the laboratory at that station, for their kindness and cooperation in making the identification of species of fleas. To them belongs the credit for the truly scientific part of the survey.

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COURT DECISION RELATING TO PUBLIC HEALTH

Law prohibiting the adulteration of coffee upheld.—(United States Circuit Court of Appeals, First Circuit; Gonzalez v. People of Porto Rico, 51 F. (2d) 61; decided June 29, 1931.) Section 1 of act 24 of the 1928 acts of Porto Rico provided as follows:

Section 1. It shall be illegal to adulterate or to mix coffee, in the grain, ground or pulverized, with any other grain or substance with the intention of selling it, or to offer or have it for sale, and it shall be equally illegal for said coffee, so adulterated or mixed, to be sold, offered or had for sale, or that it be transported or stored for the purpose of using it for human consumption, or to use it for industrial purposes, when intended for the preparation of food for human consumption.

In a prosecution for a violation of this act, it was charged that the appellant (defendant in the trial court) "unlawfully, willfully, and maliciously had and offered for sale * * * coffee roasted and ground, adulterated with another substance known as sugar." A conviction was had and this conviction was sustained by the Supreme Court of Porto Rico. On appeal to the circuit court of appeals, the contentions of appellant were (1) that the facts alleged in the information, admitted and found, did not constitute a public offense because section 1 was unconstitutional, and (2) that section 1 was invalid because in conflict with the Federal food and drugs act, which act allowed harmless adulterations provided the container or package bore a label stating the substance with which the article was adulterated and the percentage of the adulteration. The adulteration in the instant case was not injurious to health and the package bore a label stating that the coffee was mixed with 4½ per cent of sugar.

The statement by the Supreme Court of Porto Rico as to the object of the law was quoted by the circuit court of appeals as follows:

The purpose of the law was to protect the public against fraud and deceit by discouraging the admixture of cheaper or inferior grain or other substance, whether wholesome or unwholesome, which would increase the weight and impair the quality of coffee as such.

The appellate court then proceeded to hold that the legislature had acted within its constitutional powers in enacting the statute.

With respect to the appellant's second contention, the circuit court of appeals took the view that the court below had not erred "in holding that the national food and drugs act did 'not forbid the enactment of any local law prohibiting the manufacture of, or traffic in, food or other things'; and that there was 'no conflict between that statute and the law now under consideration.'"

DEATHS DURING WEEK ENDED OCTOBER 3, 1931

Summary of information received by telegraph from industrial insurance companies for the week ended October 3, 1931, and corresponding week of 1930. (From the Weekly Health Index, issued by the Bureau of the Census, Department of Commerce)

Policies in force	Week ended Oct. 3, 1931 74,736,758	Corresponding week, 1930 75,450,406
Number of death claims		12,460
Death claims per 1,000 policies in force, annual rate		8.6
Death claims per 1,000 policies, first 40 weeks of		3-47
year, annual rate	9. 8	9.7

Deaths 1 from all causes in certain large cities of the United States during the week ended October 3, 1931, infant mortality, annual death rate, and comparison with corresponding week of 1930. (From the Weekly Health Index, issued by the Bureau of the Census, Department of Commerce)

[The rates published in this summary are based upon midyear population estimates derived from the 1930 census]

	We	ek ended	1 Oct. 3,	1931		ponding , 1930		ate ² for rst 40 eks
City	Total deaths	Death rate 1	Deaths under 1 year	Infant mor- tality rate 3	Death rate ²	Deaths under 1 year	1931	1930
Total (82 cities)	6, 593	9. 6	584	4 46	10.0	660	12.0	12.0
Akron	37	7. 5 10. 9	6	59	9.4	5	7.8	7.1
Albany 5	27 72	10.9 13.5	1 9	20 92	9. 8 11. 7	3 9	13. 8 15. 2	14. 9 15. 7
Atlanta	39	10. 0	5	79	11. 6	4	10. 2	10.
Colored	33	10.8	4	115	10.8	5	(f) 14.5	(9)
Baltimore 5	168	10.8	15	51	10.8	21	14.5	14. (
White	118 50	(4)	8 7	35 109	(6)	14	(6)	(6)
ColoredBirmingham	60	(6)	2	20	12.2	3	13.6	(6)
White	35		1	17		0		
Colored	25	(6) 13. 7	1	24 57	11.6	3	(6) 14.3	(9)
Boston	207	13.7	20	57	11.6	.24	14.3	14.1
Bridgeport	19 125	6.7	13	33 53	7. 8 10. 8	11	11.1 13.2	11.1
Buffalo Cambridge	22	10. 1	0	0	14.7	6	12.2	13.0
Camden	22	9.6	Ö	ő	10. 5	1	14.4	13. 6
Canton	20	9.8	1	23	6.4	1	10.2	13. 6 10. 0
Chicago 5	569	8.6	43	38	8.9	50	10.8	10. 8
Cincinnati	118	13.5	16	96	13.4	21	16.1	15. 6
Cleveland Columbus	176	10. 1 11. 1	17	49 39	9.4	19	11. 3 13. 7	11. 2 15. 6
Dallas	63	6.9	2	09	7.3	5	11.2	11.5
White	21	0.0	ī			4		
White Colored Dayton	15	(6)	1		9.5	1	(6)	(8)
Dayton	41	10.3	4	56	9.5	4	11.9	10. 6
Denver	63	11.3	11	106	13. 4	7	14.0	14.8
Des Moines	29 217	10. 5 6. 8	3 27	53	11.3 8.7	35	11. 1 8. 3	11.8 9.4
Duluth	18	9. 2	i	43 25	10.8	2	11.4	11.3
El Paso	14	7.0	6		14.7	11	15.8	17.5
Erie Fall River 7 8	15	6.6	0	0	11.7	3	10.6	11.3
Fall River 7 8	24	10.9	1	23	7.7	0	11.2	12.0
FlintFort Worth	12	3.8 7.5	4	. 51	8. 3 9. 8	6	7.0	9.3
White	24 16	1.0	1		v. 8	4	10.9	11.1
Colored	8	(0)	1 0		(8)	ő	(0)	(8)
Grand Rapids	24	7.3	4	59	6.2	i	9.1	(6)
Houston	59	9. 9	5		11.8	6	11.3	12.2
White	41		3		(4)	5	(6)	(4)
Indianapolis	18 91	12.8	2 4	33	14.0	1 3	14.0	(6) 14.8
White	70		3	28		3	24.0	44.0
Colored	21	(6) 8. 2	1	67	(°) 8.7	ő	(6)	(8)
Jersey City	50	8.2	6	53	8.7	0 7	11.6	11.3
Kansas City, Kans	19	8.1	. 3	62	12.4	2	12.7	11.7
White	19	(6)	3 0	74	(6)	1	(6)	(8)
Kansas City, Mo	81	10.3	5	38	11.0	3 !	13. 2	13.3
Knoxville	16	7.6	2	43	8.8	2	12.6	13.7
White	11		2 0	48		2 0		*******
Colored	5	9. 2		0	10.5		9.8	9,9
Long Beach Los Angeles	27 216	8.5	9	26	8.1	18	10.7	11.0
Louisville.	59	10.0	5	43	11.2	6	14.4	13.6
White	44		4	39		6		
Colored	15	(°) 13. 5	1	66	11.9	0	12.7	(0)
Lowell	26		4	102	11.9	8	12.7	13.4
Lynn Memphis	16	8.1	1	26	9.7	8	9.6	10.5 17.2
White.	78 41	15.7	13	138	10.3	8	16.7	11.2
Colored	37	(0)	4	116	(6)	5	(6)	(8)
Miami	22	10. 2	1	25	8.9	i	11.9	11.1
. White	16		0	0		o l		

Deaths from all causes in certain large cities of the United States during the week ended October 3, 1931, infant mortality, annual death rate, and comparison with corresponding week of 1930. (From the Weekly Health Index, issued by the Bureau of the Census, Department of Commerce)—Continued.

1000	We	ek ended	1 Oct. 3,	1931		ponding , 1930	Death r	rst 40
City	Total deaths	Death rate	Deaths under 1 year	Infant mor- tality rate	Death rate	Deaths under 1 year	1931	1930
Milwaukee	97 78 41	8.6 8.6 13.7	5 6 8	22 39 119	8. 8 9. 5 15. 9	11 3 10	9. 4 11. 3 17. 0	9. 7 10. 7 16. 6
White	20 21 28 29 102	(5) 13.0 9.3 11.4	4 4 2 0 11	236 53 0	(*) 11. 1 11. 9 15. 2	6 0 7 15	(6) 12.2 12.4 17.0	(*) 11. (*) 12. 8 17. 4
White	65 37 1, 154 153	(°) 8.5 6.0	6 5 109 11 50	50 81 46 25 53	(*) 8.4 6.0 7.7	8 7 99 10	(*) 11. 3 8. 3 10. 4	(6) 10.9 8.6 9.9
Brooklyn Borough Manhattan Borough Queens Borough Richmond Borough Newark, N. J.	409 422 131 39 70	8.1 12.1 5.9 12.4 8.2	39 7 2 5	66 19 36 26	11. 8 6. 3 13. 7 10. 8	32 13 3 10	17. 0 7. 3 13. 9 11. 7	16. 1 7. 1 14. 4 12. 1
Oakland Oklahoma City Omaha Paterson	64 23 52 34 25	11. 4 6. 1 12. 5 12. 8 12. 0	4 4 8 8	51 55 45 138 105	12. 2 7. 8 16. 0 13. 5 7. 4	4 2 3 6	10 6 10 9 13. 9 13. 4 12. 6	11. 0 10. 8 13. 7 12. 4
Philadelphia Pittsburgh Portland, Oreg Providence	363 125 72 56	9.6 9.6 12.2 11.5	26 17 1 8	38 59 12 74	10.6 10.6 10.0 13.4	36 18 2 7	13. 2 14. 6 11. 6 12. 8	12.7 13.8 12.1 13.1
Richmond White Colored Rochester St. Louis	41 25 16 66 157	(6) 10. 4 9. 9	3 1 2 9	44 22 87 82 47	11. 1 (6) 9. 2 9. 9	1 1 1 2 12	(6) 12.0 15.3	(°) 11. ! 14. 2
St. Paul Salt Lake City Sen Antonio San Diego	41 26 68 48	7. 7 9. 5 14. 8 16. 0	5 3 9 2	52 45 41	9, 2 7, 8 8, 7 12, 2	2 0 4 3	10.8 12.2 14.6 13.7	10. 1 12. 2 16. 7 14. 4
San Francisco Schemectady Seattle Somerville South Bend	164 15 79 14	13. 2 8. 1 11. 1 6. 9 8. 7	3 0 4 1	20 0 38 37 50	12. 1 12. 5 10. 1 9. 5 9. 4	6 3 2 1 5	13. 1 10. 5 11. 4 9. 0 8. 1	13. 1 11. 4 10. 9 9. 8 8. 9
Spokane Springfield, Mass Springfield, Mass Tacoma	20 35 29 20	9. 0 12. 0 7. 1 9. 7	2 1 1 0 2 5	26 15 0 51	13. 1 11. 8 10. 2 10. 2	2 2 3 1	12.4 11.8 11.6 12.0	12. 4 12. 2 11. 6 12. 5
Toledo Trenton Utica Washington, D. C	79 19 26 128	13. 9 8. 0 13. 2 13. 5	1 1 13	46 17 26 72 49	12. 5 14. 8 11. 3 12. 1	12 4 1 17	12. 0 16. 5 14. 1 15. 9	12.7 16.6 14.8 15.1
White. Colored Waterbury Wilmington, Del. 7.	83 45 8 22 38	(*) 4. 1 10. 8 10. 0	6 7 9 3 5	120 0 65 69	(6) 7. 3 11. 7 11. 5	8 9 1 6 3	(5) 9.7 14.0 12.1	(8) 9. 8 14. 5 12. 8
YonkersYoungstown	13 24	4. 9 7. 2	0	56	8.9 10.4	5	8.6 10.2	8. 1 10. 3

Deaths of nonresidents are included. Stillbirths are excluded.
These rates represent annual rates per 1,000 population, as estimated for 1931 and 1930 by the arithmetical method.

¹ Deaths under 1 year of age per 1,000 live births. Cities left blank are not in the registration area for births.

births.

⁴ Data for 77 cities.

⁵ Deaths for week ended Friday.

⁶ For the cities for which deaths are shown by color, the percentage of colored population in 1920 was as follows: Atlanta, 31; Baltimore, 15; Birmingham, 39; Dallas, 15; Fort Worth, 14; Houston, 25; Indianapolis, 11; Kansas City, Kan., 14; Knoxville, 15; Louisville, 17; Memphis, 3s; Miami, 31; Nashville, 30; New Orleans, 26; Richmond, 32; and Washington, D. C., 25.

⁷ Population Apr. 1, 1930; decreased 1920 to 1930, no estimate made.

PREVALENCE OF DISEASE

No health department, State or local, can effectively prevent or control disease without knowledge of when, where, and under what conditions cases are occurring

UNITED STATES

CURRENT WEEKLY STATE REPORTS

These reports are preliminary, and the figures are subject to change when later returns are received by the State health officers

Reports for Weeks Ended October 10, 1931, and October 11, 1930

Cases of certain communicable diseases reported by telegraph by State health officers for weeks ended October 10, 1931, and October 11, 1930

	Diph	theria	Influenza		Me	osles	Meningococcu meningitis	
Division and State	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930
New England States:								
Maine	4	2	1	7	46		0	0
New Hampshire	i	10	-		1		0	i
Vermont	-	4			1		0	0
Massachusetts	56	47	4	6	22	28	1	1
Rhode Island	2	25	1		53	1	Ö	-
Connecticut.	6	5	1	2	11	9	1	i
Middle Atlantic States:				-			-	
New York	03	75	12	17	58	52	5	10
New Jersey	15	63	4	5	2	34	4	2
Pennsylvania	81	90			118	52	7	2
East North Central States:	OA.	00			220	0.0		
Ohio	111	44	7	8	9	10	1	9
Indiana	36	41		4	7	2	i	3
Illinois	79	131	62	24	8	17	4	3
Michigan	29	47	2	1	25	36	4	10
Wisconsin	16	24	14	25	12	67	2	3
West North Central States:	10	21	1.4	20	14	0.	-	
Minnesota	15	13	1		2	7	3	1
Iowa	6	9			î	4	1	i
Missouri	73	43	1	2	1	32	9	3
North Dakota	5	2	1	2	18	8	1	0
	17	13			9	1	1	1
South Dakota		13			9	4	1	0
Nebraska	17				10		0	1
Kansas	19	18	3	1	10	i	1	

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¹ New York City only.

Cases of certain communicable diseases reported by telegraph by State health officers for weeks ended October 10, 1931, and October 11, 1930—Continued

	Dip	htheria	Infl	uenza	Me	easles		goeoecus ingitis
Division and State	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930
South Atlantic States:		-					0	
Delaware	68 10	32 22		5	3	5 2	0 2	1
West Virginia	55	28	19	8	9	15	3	
North Carolina	199	173	2	10	14	3	2	
South Carolina	32 32	58 21	154 11	251 24	4	10	1 0	1 8
Florida 3	18	13	1	24	16	1	0	1
East South Central States:								
Kentucky	175	9				37	2 2 4	8
	171	60	8	16 20	11	28	2	8
Alabama. Mississippi. West South Central States:	138	38		20	11	20	0	0
West South Central States:						1		
Arkansas	44	12		15	3	1	0	3
Oklahoma •	22 99	14	3	1	2	1	0	0
Texas	35	66 25	12	1 12	2	5 2	0	0
Mountain States:		-0						1
Montana	1	6			10		0	1 0 0 1 1 1 4
Idaho Wyoming Colorado New Mexico	3				2	6	0	0
Colorado	11	1 7			3	27	0	1
New Mexico	9	11				5	Ô	i
Arizona	6	9	7	1	1	9	2	4
Utah *Pacific States:	1	2		4	1	1	0	0
Washington	6	22			7	2	0	3
Oregon	1	2	22 73	6	6	21	1	1 2
California	61	55	73	26	71	62	3	2
	Polion	nyelitis	Scarle	t fever	Sma	lipox	Typho	id fever
Division and State	Week ended	Week ended	Week ended	Week ended	Week ended	Week	Week ended	Week ended
	Oct. 10, 1931	Oct. 11, 1930	Oct. 10, 1931	Oct. 11, 1930	Oct. 10, 1931	Oet. 11, 1930	Oct. 10, 1931	Oct. 11, 1930
New England States:								
Maine New Hampshire	8	16	9 5	6	0	0	3	5
Vermont	6	2 0	4	2 2 87	1	0	1 2	0
Vermont	72	53	151	87	1 0	0	12	9
Khode Island	. 5	2	7	0	0	0	0	11
Connecticut	45	10	9	16	0	0	5	11
Middle Atlantic States: New York New Jersey	239	51	184	111	0	0	35	35
New Jersey	50	9	54	49 141	0	0	12	11
Pennsylvania. East North Central States:	40	9	187	141	0	0	69	139
Ohio	8	26	178	174	0	3	57	49
indiana	5	14	48	81	3		12	15
Illinois	61	27	178	193	16	8 9 2	51	28
Michigan	74 49	15 16	102	119 62	1	0	20	28 83 3
Wisconsin. West North Central States:	49	10	22	62	1	0	9	
Minnesoth	58 13	13	36	33	0	3	3 5	1
lowa	13	21	31	39	5	15	8	1 2 24
Missouri North Dakota	7	27	107	42 12	8 5	10	15	24
South Dakota	ő	24	7	8	1 2	5	8	1
Nobeceles	1	15	18	14	i	9	1	
Nebraska	4 1	57	46				13	13

Week ended Friday.
 Typhus fever, 1931, 11 cases: 1 case in Maryland, 2 cases in South Carolina, 6 cases in Georgia, and 2 cases in Florida.
 Figures for 1931 are exclusive of Oklahoma City and Tulsa.

Cases of certain communicable diseases reported by telegraph by State health officers for weeks ended October 10, 1931, and October 11, 1930—Continued

	Polion	yelitis	Scarle	t fever	Sma	llpox	Typho	id fever
Division and State	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11, 1930	Week ended Oct. 10, 1931	Week ended Oct. 11 1930
South Atlantic States: Delaware Maryland **	1 5	0 3	5 61	4 33	0	0	2 33	10
District of ColumbiaVirginia	3	1	15	10	0	0	9	
Virginia West Virginia		3	43	48	0	1	79	58
North Carolina	7	1	111	109	3	ô	23	23
South Carolina 3	0	i	9	22	0	0	22	40
Georgia 1	0	3	34	32	2	0	28	37
Florida 3	0	0	0	6	0	1	3	1
East South Central States:		2.1						
Kentucky	1	3	68	27	0	5	68	30
Tennessee	3	5	63	54	1	2	30	41
Alabama	0	3	66	66	0	1	33	- 15
Mississippi	0	2	40	26	1	1	27	10
West South Central States:	- 1						0.0	
Arkansas	0	4	23	7	1	5	19	45
Louisiana	1	- 3	17	9	2	0	40	21
Oklahoma 4	0	8	34	47	1	2	56	37 11
Texas.	0	10	39	11	5	11	36	11
Mountain States: Montana	-		10	26	0			
	7	1 0	10	6	8	0	9	0
Wyoming	ő	2	5	4	0	0	1	0
Colorado	1	- 1	12	8	0	1	i	19
New Mexico.	4	9	7	9	ő	i	14	19
Arizona	i	î	i	3	ő	ô	2	13
Utah 2	ô	ô	6	11	1	0	4	1
Pacific States:		-	-		-			
Washington	10	1	26	40	5	10	4	12
Oregon	0	0	8	11	1	0	3	3
California	6	57	67	75	9	22	15	13

SUMMARY OF MONTHLY REPORTS FROM STATES

The following summary of cases reported monthly by States is published weekly and covers only those States from which reports are received during the current week:

State	Menin- gocoe- cus menin- gitis	Diph- theria	Influ- enza	Ma- laria	Mea- sles	Pel- lagra	Polio- myelitis	Scarlet fever	Small- pox	Ty- phoid fever
September, 1931										
Alabama	5	299	13 13	373	26	110	10	156	3	127
Arizona	- 4	16	13	1	10	2	2	15 26 112	0	27
Connecticut	2	20	7	1	17		458	26	0	23
Indiana	5	56	46		28		12	112	31 17	66
Iowa	4	33			9		34	50	17	14
Maine	2	13	1		36		20	18	0	18 97
Michigan	24	13 73	4	1	59		20 577	285		97
North Dakota	4	5			7		11	15	6	17
Porto Rico		44	73	2,840	11		1		0	11

September, 1931		Conjunctivitis, infectious:	8505
Chicken pox:	ases	Connecticut	. 1
Alabama	. 21	Maine	1
Arizona	. 5	Dengue:	
Connecticut	. 32	Alabama	1
Indiana	. 21	Dysentery:	
Iowa	. 17	Arizona	5
Maine	. 11	Connecticut (bacıllary)	1
Michigan	. 85	Porto Rico	45
North Dakota	. 9	Filariasis:	
Porto Rico	. 3	Porto Rico.	. 8

Week ended Friday.
 Typhus fever, 1931, 11 cases: 1 case in Maryland, 2 cases in South Carolina, 6 cases in Georgia, and 2 cases in Florida.
 Figures for 1931 are exclusive of Oklahoma City and Tulsa.

German measles:	Cases	Tetanus:	ases
Arizona	. 1	Connecticut	. 2
Connecticut	. 9	Maine	. 1
Iowa	. 2	Porto Rico	. 5
Maine	. 5	Tetanus, infantile:	
Lead poisoning:		Porto Rico	. 12
Connecticut	. 2	Trachema:	
Leprosy:		Arizona	. 29
Porto Rico	. 1	North Daketa	2
Lethargic encephalitis:		Porto Rico	6
Alabama	. 4	Trichinesis:	
Connecticut	. 3	Connecticut	1
Michigan	. 8	Typhus fever:	
Mumps:		Alabama	5
Alabama	. 13	Maine	18
Arizona	. 7	Undulant fever:	
Connecticut		Alabema	1
Indiana		Arizona	1
Iowa	. 19	Indiana	1
Maine		Iowa	2
Michigan		Maine	2
North Dakota		Michigan	
Porto Rico		North Dakota	1
Ophthalmia neonatorum:		Vincent's angina:	
Arizona	. 1	Maine	5
Connecticut	1	North Dakota	48
Porto Rico		Whooping cough:	
Paratyphoid fever:		Alabama	81
Connecticut	4	Arizona	3
Porto Rico	1	Connecticut	258
Puerperal septicemia:		Indiana	120
Porto Rico	6	Iowa	92
Rabies in animals:		Maine	35
Connecticut	4	Michigan	848
Rabies in man:		North Dakota	83
Alabama	1	Porto Rico	119
Septic sore throat:			
Connecticut	5		
Iowa	-		
Miebigan			

Cases of Certain Communicable Diseases Reported for the Month of May, 1931, by State Health Officers

State	Chick- en pox	Diph- theria	Measles	Mumps	Scarlet fever	Small- pox	Tuber- cu- losis	Ty- phoid and para- typhoid fever	Whoop- ing cough
Maine	1, 138	23 6 2 152 20 39	68 69 2, 299 505 2, 414	246 95 644 257 276	145 11 22 1, 542 226 200	0 0 18 0 0	75 19 491 61 105	6 0 2 21 2 9	103 42 626 36 172
New York New Jersey Pennsylvania	1, 827	536 166 297	12, 992 4, 190 16, 967	1, 744 296 1, 778	3, 650 1, 160 2, 600	32 6 0	1, 755 489 627	72 15 45	1, 920 933 853
Ohio. Indiana. Iliinois Michigan. Wisconsin	1, 916 364 1, 402 1, 439	134 81 481 137 65	5, 027 4, 501 8, 350 787 3, 442	2, 511 205 1, 060 812 4, 544	1, 824 913 2, 149 1, 697 624	192 541 265 81 50	749 931 879 543 158	39 11 25 16 4	481 344 815 1, 067 609
Minnesota Iowa Missouri North Dakota Bouth Dakota Nebraska Kansas	1, 032 186 305 131	52 24 160 30 41 26 46	897 271 2, 419 302 186 49 497	105 198 113 10 655 557	344 237 1, 340 145 52 198 170	33 274 212 22 29 233 284	268 34 262 16 20 20 136	8 1 35 5 3 3 10	256 108 300 51 43 111 176

Cases of Certain Communicable Diseases Reported for the Month of May, 1931, by State Health Officers—Continued

State	Chick- en pox	Diph- theria	Measles	Mumps	Scarlet fever	Small- pox	Tuber- cu- losis	Ty- phoid and para- typhoid fever	Whoop ing cough
Delaware	18	2	539	34	64	0	20	3	12
Maryland	330	47	4, 589	313	287	0	1 221	24	258
District of Columbia	86	37	1, 222		76	0	97	3	35
Virginia.	642	67	3, 605		159	13	217	39	461
West Virginia	284	33	646		190	27	75	27	274
North Carolina	445	60	3, 996		169	13		17	846
South Carolina	392	82	674	152	28	6	183	47	304
Georgia	179	31	823	175	276	44	127	48	172
Florida	161	19	764	43	20	5	47	12	71
Kentucky 1							******		
Tennessee	188	42	1,704	164	414	100	263	32	291
Alabama	148	44	1, 110	102	100	56	582	38	92
Mississippi	694	33	260	331	78	184	155	45	450
Arkansas	109	10	212	67	50	106	1 27	27	68
Louisiana	108	74	22	8	84	74	1 163	49	19
Oklahoma 1	208	42	-183	31	108	280	59	24	60
Texas		97			147			39	
Montana	167	7	70	80	80	4	48	5	97
Idaho	39	9	22	16	52	10	1	6	109
Wyoming	35		6	72	45	2		0	32
Colorado	249	21	894	193	136	30	69	3	324
New Mexico	85	14	424	65	25	8	45	8	
Arizona Utah ³	26	13	215	15	11	0	93	10	32
Nevada	22		88	4	1	0	1	0	
Washington	578	36	1, 028	264	144	104	227	28	541
Oregon	222	31	424	255	74	90	59	- 8	78
California	1,710	304	4, 780	1, 145	554	93	834	45	1, 166

¹ Pulmonary.

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Case Rates per 100,000 Population (Annual Basis) for the Month of May, 1931

State	Chiek- en pox	Diph- theria	Mensles	Mumps	Scarlet fever	Small- pox	Tuber- cu- losis	Ty- phoid and para- typhoid fever	Whoop- ing cough
Maine New Hampshire Vermont Massachusetts Rhode Island Connecticut	244 415 312 128 282	34 15 7 42 34 28	100 225 630 852 1,739	362 310 176 434 199	213 28 72 422 381 144	0 0 59 0 0	110 62 135 103 76	9 0 7 6 3 6	137 171 61 124
New York New Jersey Pennsylvania	249 518 297	49 47 36	1, 190 1, 189 2, 051	160 84 215	334 329 314	3 2 0	161 139 76	7 4 5	176 265 103
Ohlo	334 131 212 340 768	23 29 73 32 26	876 1, 618 1, 265 186 1, 362	438 74 161 192 1,798	318 328 326 401 247	33 194 40 19 20	131 335 133 128 78	7 4 4 4 2	84 124 123 257 241
Minnesota Iowa Missouri North Dakota South Dakota Nebraska Kansas	470 88 98 225 121 279 208	24 11 52 52 69 22 29	408 129 779 519 313 42 309	50 64 194 17 556 346	157 113 431 249 88 168 106	15 130 68 38 99 198 176	131 16 91 28 34 17 85	4 0 11 9 5 3 6	117 51 97 88 79 94 109
Delaware	88 235 205 310 190 161 264 72 124	10 33 88 32 22 22 25 55 13 15	2, 641 3, 267 2, 919 1, 743 432 1, 450 455 333 588	167 223 103 71 33	314 204 182 77 127 61 19 112 18	0 0 0 6 18 5 4 18	98 1 157 232 105 50 123 51 36	15 17 7 19 18 6 32 19	04 184 84 223 183 307 206 70 58

¹ Pulmonary.

² Reports received weekly. ³ Exclusive of Oklahoma City and Tulsa.

Case Rates per 100,000 Population (Annual Basis) for the Month of May, 1931—Continued

State	Chick- en pox	Diph- theria	Measles	Mumps	Scarlet fever	Small- pox	Tuber- cu- losis	Ty- phoid and para- typhoid fever	Whoop- ing cough
Kentucky 1									
Tennessee	84	19	757	73	184	44	117	14	129
Alabama	65	19	487	45	44	25	255	17	40
Mississippi	401	19	150	191	45.	106	90	26	260
Arkansas	69	6	134	42	32	67	1 17	17	43
Louisiana	59	41	12	4	46	41	1 90	27	10 34
Oklahoma	117	24	103	17	61	157	83	13	34
Texas		19	******		29			8	
Montana	366	15	153	175	175	9	105	11	212
Idaho	103	24	58	42	137	26	3	16	287
Wyoming	180		31	870	231	10		0	164
Colorado	280	24	1,005	217	. 153	34	78	3	364
New Mexico	232	38	1, 158	178	68	. 22	123	22	
Arizona	68	34	565	39	29	0	244	22 26	84
Utah 1					*******	******			
Nevada	279		1, 117	81	13	0	13	0	
Washington	428	27	762	196	107	77	168	21	401
Oregon	268	37	512	308	89	109	71	10	91
California	338	60	946	227	110	18	165	9	231

¹ Pulmonary.

GENERAL CURRENT SUMMARY AND WEEKLY REPORTS FROM CITIES

The 96 cities reporting cases used in the following table are situated in all parts of the country and have an estimated aggregate population of more than 33,315,000. The estimated population of the 91 cities reporting deaths is more than 31,935,000. The estimated expectancy is based on the experience of the last nine years, excluding epidemics.

Weeks ended October 3, 1931, and October 4, 1930

	1931	1930	Estimated expectancy
Cases reported	18		100
Diphtheria:			
46 States	1,726	1, 228	
96 cities	356	377	600
Measles:			- Indian
45 States	451	644	
96 cities	116	116	
Meningococcus meningitis:	4		
46 States	20	77	
96 cities	20	32	
Poliom velitis: 46 States	956	649	
Scarlet fever:			
46 States	1,607	1,686	
96 cities	419	450	463
Smallpox:		-	-
46 States	105	175	
96 cities	1	5	2
Typhoid fever:			Car Charles
46 States	1,049	933	
	135	124	143
96 cities	100	103	***
Deaths reported	1000		ALC: U.S.
a leading to the second		-	
Influenza and pneumonia: 91 cities	842	366	
Smallpox: 91 cities	0	0	

^{*} Reports received weekly.

¹ Exclusive of Oklahoma City and Tulsa.

City reports for week ended October 8, 1931

The "estimated expectancy" given for diphtheria, poliomyelitis, scarlet fever, smallpox, and typhoid fever is the result of an attempt to ascertain from previous occurrence the number of cases of the disease under consideration that may be expected to occur during a certain week in the absence of epidemics. It is based on reports to the Public Health Service during the past nine years. It is in most instances the median number of cases reported in the corresponding weeks of the preceding years. When the reports include several epidemics, or when for other reasons the median is unsatisfactory, the epidemic periods are excluded, and the estimated expectancy is the mean number of cases reported for the week during nonepidemic years.

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If the reports have not been received for the full nine years, data are used for as many years as possible, but no year earlier than 1922 is included. In obtaining the estimated expectancy, the figures are smoothed when necessary to avoid abrupt deviation from the usual trend. For some of the diseases given in the table the available data were not sufficient to make it practicable to compute the estimated expectancy.

Division, State, and city	Chicken pox, eases reported	Diphtheria		Influenza				
		Cases, estimated expectancy	Cases reported	Cases reported	Deaths reported	Measles, eases re- ported	Mumps, cases re- ported	Pneu- monia, deaths reported
NEW ENGLAND								
Maine:								
Portland	. 0	0	1	L	0	1	1	
New Hampshire:					110			
Concord	0	0	0		0	0	. 0	
Manchester	8	0	0		0	0	0	
NashuaVermont:	0	0	0		0	0		
Barre		0	0		0			
Massachusetta:	-							,
Boston	5	16	12	3	1 0	2	2 0	8
Fall River	1	3	2		0	4	0	1
Springfield	1	1			0	0	3	
Worcester Rhode Island:		4	0		0	0	11	
Pawtucket		1	0			0		
Providence	. 0	1	. 5		ŏ	3	1	
Connecticut:					1 1 1 1 1 1			
Bridgeport	1 0	3 2	0		0	0	0	
Hartford	0	2	1 0		6	0	0	9
New Haven						0	•	1
					4			
New York:			1					1111
Buffalo	2	9	2 41		0	0	0	10
New York Rochester	13	88	41	10	3	10	17	93
Syracuse	1 0	2	1 0		0	3	2	,
New Jersey:			U					
Camden	0	3	0		0	0		0
Newark	1 0	10	2	2	0	0		3
Trenton	0	. 1	0		0	0	3	1
Pennsylvania: Philadelphia	4	-	1		3	1		10
Pittsburgh	3	33 13	8			12	8	11
Reading	ĭ	1	O		o	0	0	1
EAST NORTH CENTRAL			1				- 2	A.C.
Ohio:							5.00	100
Cincinnati	0					. 0	. 0	5
Cleveland	8	31	4	2	1	3	12	10
Columbus	i	4	15		ő	i	1	i
Toledo	0	. 0	3		0	4	Ö	
indiana:			1			1000		tole ?
Fort Wayne Indianapolis	0	1	3		0	0	0	0
South Bend	0	8	1		1	1	1	
Terre Haute	1	ő	i		0	ő	o	i
Minois:	1 0 0	0					U	2 14
Chicago	8	67	39	1	1	6	10	26
Springfield	1	0			0			1

City reports for week ended October 3, 1931-Continued

Division, State, and city	Chicken pox, cases reported	Diphtheria		Influenza			- 30	
		Cases, estimated expect- ancy	Cases reported	Cases reported	Deaths reported	Measles, cases re- ported	Mumps, cases re- ported	Pneu- monia, deaths reported
EAST NORTH CEN- TRAL—continued				361			Aller d	
Michigan:		-					100	etil .
DetroitFlint	1 0	39	7 0	2	0	0	5 3	2
Grand Rapids	0	1	- 0		Ŏ	1	0	0
Wisconsin: Kenosha	0	0	0	1.80	0	0	4	
Madison	ő	2 7	0			i	5	
Milwaukee	9	7	0		0	1 5 0	10	3
Racine Superior	0	0	0		0	1	2	3
WEST NORTH CENTRAL		4 1			10		- 1	
Minnesota: Duluth	,	0	0	and the	0	0	0	1
Minneapolis	17	21	7		0 2	1 0	18	1
St. Paul	3	9	3		0	0	0	2
Davenport	. 0	1	0			0	0	
Des Moines	0	2	1			0	0	
Sioux City Waterloo	0	1 0	3 0			0	1 0	
Missouri:	0							
Kansas City	3	4	2		0	0	2	3
St. Joseph St. Louis	0 2	0 24	3 14		0	0	0	2
North Dakota:							243	
Fargo	0	1	0		0	. 0	0	1
Grand Forks South Dakota:	0	0	0			. 0	0	
Aberdean	16	0	0			. 8	0	
Nebraska:			***					
Omaha Kansas:	0	8	12		0	0	- 1	
Topeka	0 2	1 2	1 0		0	0	3	1
SOUTH ATLANTIC			-					1000
Delaware:							46.0	193 -
Wilmington	0	1	0		0	0	0	
Maryland: Baltimore	4	17	13	1	0	1	6	
Cumberland	0	0	0	-	0	0 0	0	· Ó
Frederick	0	0	0		0	0	0	0
District of Columbia: Washington	1	11	9		0	0	0	
Virginia:	1	**					27 1	(r)-1 - C
Lynchburg	0	3			000	. 0	0	1
Norfolk Richmond	0	17	0 14		0	. 0	1 0	
Roanoke	0 0	3	10		ŏ	. 0	0	1
West Virginia:								
Charleston	0	1 0	0	1	0	0	0	0
NORTH Carolina: 1								379
Raleigh	0	3	1		0	0	0	0
Wilmington Winston-Salem	0	1	13		0	0	0 2	
South Carolina:		100	20				10000	1,500
Charleston	0	1	1	7	0	0	0	2
Columbia Greenville	0	1 2	0 2		0	0	0	8 0
Georgia:		-						41 11 11
Atlanta	0	7	6	4	0	0	0	2
Brunswick Savannah	0	0	0	3	0	0	0	0
Florida:	0	1	2		0	0	0	
Miami	0	2	0		0	17	1	1
Tampa	0	11	1		0	0	0	0

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City reports for week ended October 3, 1931-Continued

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		Diph	theria	Infly	uenza			9
Division, State, and city	Chicken pox, cases reported	s Cases,	Cases reported	Cases reported	Deaths reported	Measles, cases re- ported	Mumps, cases re- ported	doothe.
EAST SOUTH CENTRAL								
Kentucky: Covington Tennessee:	. 0	1	1		. 0			
Memphis Nashville	0		15		0	0	0	
Alabama: Birmingham Mobile	0	1	3 4		1 0	0	0	0
Montgomery	0	3	ō			5	2	
WEST SOUTH CENTRAL Arkansas:								- 3
Fort Smith Little Rock	8	8	- 0		0	0	0	
Louisiana: New Orleans Shreveport	. 0	8	7 3		0		8	
Oklahoma: Muskegee Oklahoma City	0	0 2	13		0	0	0	9
Tulsa Texas: Dallas	0	10	38		0	0	0	,
Fort Worth Galveston	8	10 2 0 8	5 0 8		0	0	0	0
Houston San Antonio	0	2	*		0		ō	i
MOUNTAIN Montana:								
Montana: Billings Great Falls Helena	. 0	0	0		0 0	0 1	0	
MissoulaIdaho:	0	9	0			0	0	1
Boise Colorado: Denver	8	1 8	7		0	0	0	,
Pueble New Mexico:	0	0	0		0	0	o o	
Albuquerque Arizona: Phoenix	0	1	0		0	0	0	1
Utah: Salt Lake City Nevada:		2	2		0	0	0	,
Reno	0	0	0		0	0	0	,
PACIFIC Washington:								
Seattle Spokane Tacoma	12	1	0			8	3	
Tacoma Oregon: Portland	12	3 8	0		0	0 2	1	
SalemCalifornia:	0	0	0		0	0	ŏ	
Bacramento San Francisco	0 14	21 2 10	19 0 2	14	0	12 10 13	0	3

	Scarle	t fever		Smallpo	x	Tuber-	T	rphoid f	ever	Whoop	
Division, State, and city	Cases, esti- mated expect- ancy	Cases re- ported	Cases, esti- mated expect- ancy	Cases re- ported	Deaths re- ported	culo- sis, deaths re-	mated	Cases re- ported	Deaths re- ported	ing cough, cases re- ported	Deaths, all causes
NEW ENGLAND		1/3									
Maine:									-		
Portland New Hampshire:	1	1	0	0	0	0	1	0	. 0	2	14
Concord	0	2	0	0	0	0	0	0	0	0	
Manchester Nashua	0	0	0	0	0	1 0	0	0	0	0	3.5
Vermont:				-0-			0				
Barre	0	0	0	0	0	1	0	0	0	0	2
Massachusetts: Boston	24	22	0	0	0	9	3	1	0	16	207
Fall River	2 2	22 7 5	0	0	0	2	. 1	. 0	0	2	24
Springfield	2	13	. 0	0	0	2	0	0	0	2 9	35
Worcester Rhode Island:	5	13	0	. 0	0	1	0	0	0	a	35
Pawtucket	0	0	0	0	0	0	0	0	0	0	16
Providence	2	3	0	0	0	2	1	4	0	4	56
Connecticut: Bridgeport	2	1	0	0	0	0	0	0	0	0	19
Hartford	1	0	0	0	0	3	0	2 0	0	4	35
New Haven	1	1	0	0	0	3	0	0	0	3	29
MIDDLE ATLANTIC											177
New York: Buffalo	8	20	0		0	11		0	0	16	122
New York	36	25	0	0	0	79	32	19		166	1, 154
Rochester	2 2	11	0	0	0	1	2	2	3	3	63
Syracuse New Jersey:	2	4	0	0	0	7	0	0	0	16	29
Camden	2	3	0	0	0	0	1	0	0	2	22
Newark	5	8 2	0	0	0	10	2	2	0	79	74 19
Trenton Pennsylvania:	1	2	0	0	0	2	0		0	0	10
Philadelphia	27	32	0	0	0	22	11	20	0	91	363
Pittsburgh Reading	17	8	0	0	0	10	3	4	1 0	85	125 34
EAST NORTH CENTRAL			ď								
Ohio:		-						1	-67	J. J.	
Cincinnati Cleveland	8	17	0 0	0	0	12	2 3 1 1	3 1 1	0	73	118 176
Columbus		12 8	0	0	0	4	1	î	0	31	63
Toledo	5	6	0	0	0	4	1	5	0	31	79
Indiana: Fort Wayne	1	0	0	0	0	0	1	0	0	3	19
Indiananolis	6	0	0	0	Ö	9 2	2 0	0	0	0	
South Bend	2	1	0	0	0	0	0	0	0	2 0	18 15
Terre Haute	1	0	0	0	0	0	0	1	0	0	40
Chicago	45	31	0	0	0	36	6	2	1	101	589
Springfield Michigan:	1	1	0	0	0	1	0	0	0	0	18
Detroit	36	17	0	0	0	18	4	5	2	116	217
Flint	7	3	0	0	0	0	0	0	0 0	8	12 24
Grand Rapids_ Wisconsin:	6	4	0	0	0	1	1	0	0	0	24
Kenosha	0	0	0	0	0	1	0	0	0	0	4
Madison	1	0	0	0			0	0		8	
Milwaukee Racine	10	5 2	0	0	0	3	1 0	1 0	0	82	97
Superior	3	1	0	0	0	1	0	0	0	1 0	

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	Scarle	t fever		Smallpo	X	Tuber-	Т	phoid f	ever	Whoop-	
Division, State, and city	Cases, esti- mated expect- ancy	Cases re- ported	Cases, esti- mated expect- ancy	Cases re- ported	re-	culo- sis, deaths	Cases, esti- mated expect- ancy	Cases re- ported	Deaths re- ported	ing cough, cases re- ported	Deaths all causes
WEST NORTH CENTRAL											
Minnesota:											
Minneapolis St. Paul	21 11	15 4	0	0	0	3 3	0 1 1	0 1 0	0	0 6 10	18 78 48
Iowa: Davenport	0	2	0	2			0	0		0	
Des Moines	3	3	0	1			0	0		0	25
Sioux City Waterloo	2 2	4	0	0			1 0	0		2	
Missouri:		******			*******						
Kansas City	6	3	0	0	0	11 2	0	0	0	5	81
St. Joseph St. Louis	16	12	0	0	o	7	5	3	0	39	157
North Dakota:											
Grand Forks	1	0	0	0	0	0	. 0	0	0	3	
South Dakota:											
Aberdeen	1	2	0	0			0	0		8	
Nebraska: Omaha	2	4	0	1	0	3	0	1	0	1	52
Kansas:	2		0	0	0		0	1	0	0	-
Topeka Wichita	2	2 2	ő	ő	ő	0	0	ó	ő	1	26 17
SOUTH ATLANTIC				-							
Delaware:					1						
Wilmington	1	0	0	0	0	1	0	0	0	0	23
Maryland: Baltimore	8	4	0	0	0	21	8	4	0	104	168
Cumberland	0	0	0	0	0	2	1	0	0	0	12
Frederick District of Col.:	40	O	0	0	0	0	0	0	0	0	3
Washington	8	6	0	0	0	9	3	0	0	10	128
Virginia:				0				2			11
Lynchburg Norfolk	1	0	0	0	0	1 2	0	ő	1 0	0	11
Richmond	1 5	15	0	0	0	4	1	2	0	.8	44
Roanoke West Virginia:	2	1	0	0	0	1	0	1	0	0	10
Charleston	2 2	1	0	0	0	2 0	1	1 14	1	3	30 24
Wheeling	2	0	0	0	0	0	0	2	. 0	1	24
North Carolina: Raleigh	1	0	0	0	0	2	0	0	0	2	17
Wilmington	1	Ö	ő	0	0	1	0	0	0	2	12
Winston- Salem	4	1	0	0	0	1	1	0	0	2	17
South Carolina:											
Charleston	0	1	0	0	0	3	0	0	0	0	28
Greenville	0	1 0	0	0	0	ő	0	0	ő	0	
Georgia:								-1	-		-
Atlanta Brunswick	6	1 0	0	0	0	3	0	7	5	0	20 00
Savannah	0	ő	ő	Ö	0	3	i	0	0	0 2	28
Florida:	0	0	0	0	0	2	1	1	0	0	22
Miami Tampa	ő	ő	ő	ő	ő	ő	0	î	ő	3	22 16
EAST SOUTH CENTRAL	-		-								
Kentucky:									4		
Covington	1	0	0	0	0	0	0	1	0	0	12
Tennessee:											78
Memphis Nashville	3 2	5	0	0	0	9	3	2	8	14	41

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¹¹³ cases nonresidents.

	Scarle	t fever	1	Smallpo	X	Tuber-	Ty	phoid f	over	Whoop-	
Division, State, and city	Cases, esti- mated expect- ancy	Cases re- ported	Cases, esti- mated expect- ancy	Cases re- ported	Deaths re- ported	culo- sis, deaths	mated	Cases re- ported	Deaths re- ported	ing cough, cases re- ported	Deaths all causes
EAST SOUTH CENTRAL—con.		7 = 1		7	1-1						
Alabama: Birmingham Mobile Montgomery	6 1 1	3 1 3	0 0	0 0	0	1	2 0 1	2 0 0	0	3 0 4	00 21
WEST SOUTH CENTRAL				15							19
Arkansas: Fort Smith Little Rock	1 2	0	0	0	0	2	. 0	0	1	0	
Louisiana: New Orleans Shreveport	3 1	2	0	0	0	11 0	3	0	1	0 3	102
Oklahoma: Muskogee Oklahoma	0	0	0	0	0	0	0	1	0	0	
City Tulsa	3	1 2	1 0	0	0	2	2	0	0	0	23
Texas: Dallas Fort Worth Galveston Houston	3 1 1 1	7 9 0	0 0 0	0 0 0	0 0 0	1 2 0 4	2 1 0 1	1 8 0 2	0 0 0	0 0 0	36 24 8 86
San Antonio MOUNTAIN	1	0	0	0	0	9	1	0	0	0	66
Montana:						- 1		177			
Billings Great Falls Helena	0 1 0 0	0 0 0	0 0 0 1	0 0 0	0 0	0 0 0	0 0	0 0 0	0 0	1 1 0 0	
MissoulaIdaho: Boise	0	1	0	0	0	0	1	0	1	0	0.5
Colorado: Denver	6	8	0	0	0	4 0	2	1	1 0	5 0	66
Pueblo New Mexico: Albuquerque	0	0	0	0	0	0	2	6	0	0	
Arizona: Phoenix	1	1	0	0	0	2	0	0	0	0	
Utah: Salt Lake City.	2	1	. 0	0	0	2	2	0	0	1	20
Nevada: Reno	0	0	0	0	0	0	0	0	0	0	
PACIFIC			-			1 7			434	17971	-
Washington: Seattle Spokane Tacoma	7 3 1	7	0 1 1	0	0	0	2 1 1	1	0	0	20
Oregon: Portland	4	5	2 1	3	0	2 0	1	2 0	0	0	72
Salem	12	21	0	0	0	18	3		0	17	216
San Francisco.	8	5	0	0	0	13	1	i	ő	0	24 157

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	Mening	gococcus Ingitis	Letha	rgic en- alitis	Pell	lagra	Polior	nyelitis e paraly:	(infan- sis)
Division, State, and city	Cases	Deaths	Cases	Deaths	Cases	Deaths	Cases, esti- mated expect- ancy	Cases	Death
NEW ENGLAND									
Maine:									
Portland New Hampshire:	0	0	0	0	0	0	0	5	
Concord		0		0	U				
Boston Fall River Springfield	1	1	0	0	0	0	4	31	
Fall River	0	0	0	0	0	0	0	3	9
Springheid	0	0	0	0	0	0	0	10	1
Worcester Rhode Island:		0	U	0	U	0	0		,
Providence	2	1	0	0	0	0	0	4	
Connecticut:	1.3								
Bridgeport	0	0	. 0	0	. 0	0	0	12	1
Hartford	0	0	0	0	0	0	1 0	6	1
New Haven		0		0					
MIDDLE ATLANTIC						1			10
New York:									
New York	0	0	0	0	0	0	15	140	10
Rochester	0	0	0	0	0	0	1	1	
Syracuse New Jersey:	0	0	U	0	0		•		
Newark	0	0	. 0	0	0	0	1	7	. 0
Pennsylvania:		1						1	
Philadelphia Pittsburgh	0	0	0	0	0	0	1	8	0
	1	1	0	0	0	0	0	0	0
EAST NORTH CENTRAL				11 0					
Ohio:									
Toledo	0	0	0	0	0	0	0	1	1
Fort Wayne	0	0	0	0	0	0	0	2	0
Illinois 1		-				-		-	
Chicago	3	2	1	0	0	0	4	13	1
Michigan:				0				9	
Detroit	0	0	0	0	0	0	0	2	
Flint Grand Rapids	ő	0	ő	ő	ŏ	ő	1	il	ŏ
Wisconsin:			-						
Kenosha	0	0	0	0	0	0	0	2	0
Madison	1	0	0	0	0	0	0	4	9
Milwaukee	0	0	0	01	ő	0	0	3	
Superior	ŏ	0	ő	0	ő	ŏ	0	4	i
WEST NORTH CENTRAL									
Minnesota:							6		
Daleth	0	0	0	0	0	0	0	2	0
Minneapolis	0	0	0	0	0	0	2	12	
St. Paul	0	0	0	0	0	0	0	20	0
Missouri:	1	0	1	0	0		0	9	
St. Louis North Dakota:	1	0		0	0	0	0	-	
Fargo	0	0	0	0	0	0	0	1	

¹ Typhus fever, 3 cases: 1 case at Springfield, Ill., and 2 cases at Savannah, Ga.

	Mening men	gococcus ingitis		rgic en- malitis	Pell	agra	Polior	nyelitis e paralys	(Infan- sis)
Division, State, and city	Cases	Deaths	Cases	Deaths	Cases	Deaths	Cases, esti- mated expect- ancy	Cases	Deaths
SOUTH ATLANTIC									
Maryland:									
Baltimore District of Columbia:	0	0	1	0	0	0	1	1	
Washington	0	0	0	0	0	0	1	4	-
Vierinia:				1	0	0	1		
Richmond		0	0	1	U				,
Charleston	2	1	0	0	0	0	0	0	
North Carelina:		0	0		1	. 0	0	0	
Raleigh Winsten-Salem	0	0	0	0	1	1	0	0	1
South Carolina:									
Charleston	0	0	1 0	0	0	0	0	0	
Georgia		0					-		
Savannah 1	0	0	0	0	1	0	0	0	
EAST SOUTH CENTRAL	10 . 3								10.
Tennossee:									
Memphis Nashville	0	0	0	0	0	0	1	0	
Alabama:									
Birmingham	1	1	0	0	0	0	0	0	
WEST SOUTH CENTRAL									
Arkansas:									
Little Rock	0	0	0	0	0	1	0	0	,
New Orleans	1	0	0	0	0	0	0	0	
Texas:					2	2		0	
Port Worth	0	0	0	0	0	1	0	0	
Galveston	o	ŏ	ő	ő	ő	î	Ö	0	
MOUNTAIN	111							-	
New Mexico:	1.16								100
Albuquerque	0	0	0	0	0	0	0	1	
PACIFIC									N 1
Washington:					1	-	1		
Beattle.	1	0	0	0	0	0	11	0	
Tacoma.	1	0	. 0	0	0	0	1	1	0
California: Los Angeles	1	0	0	0	0	0	2	. 0	
San Francisco	i	1	1	il	1	0	î	1	

¹Typhus fever, 3 cases: 1 case at Springfield, Ill., and 2 cases at Savannah, Ga.

The following tables give the rates per 100,000 population for 98 cities for the 5-week period ended October 3, 1931, compared with those for a like period ended October 4, 1930. The population figures used in computing the rates are estimated mid-year populations for 1930 and 1931, respectively, derived from the 1930 census. The 98 cities reporting cases have an estimated aggregate population of more than 33,000,000. The 91 cities reporting deaths have more than 31,500,000 estimated population.

Summary of weekly reports from cities, August 30 to October 3, 1931.—Annual rates per 100,000 population compared with rates for the corresponding period of 1930

DIPHTHERIA CASE RATES

					Week	ended-				
1.	Sept. 5, 1931	Sept. 6, 1930	Sept. 12, 1931	Sept. 13, 1930	Sept. 19, 1931	Sept. 20, 1930	Sept. 23, 1931	Sept. 27, 1939	Oct. 3, 1931	Oct. 4, 1930
98 cities	36	40	35	44	34	46	45	56	1 56	60
New England Middle Atlantic East North Central West North Central South Atlantic East South Central West South Central Mountain Pacific	81 105 82 27	39 29 48 35 66 48 56 44 32	58 26 32 34 45 99 41 26 29	60 26 63 56 68 24 45 35 22	* 36 22 29 42 73 93 57 17 29	34 36 74 48 46 24 63 26 12	38 25 42 71 67 128 101 52 41	56 31 74 58 109 30 136 62 26	50 25 44 *88 150 140 108 78 •43	53 40 79 60 68 102 104 9 51
		MEA	SLES (CASE I	RATES					
98 cities	19	24	14	16	22	16	15	18	1 18	19
New England Middle Atlantic East North Central West North Central Fouth Atlantic East South Central West South Central Mountain Pacific	58 14 11 8 8 6 10 52 67	36 27 12 31 28 24 0 53 34	29 8 13 11 6 6 10 35 45	41 19 9 15 6 6 3 35 16	31 18 17 13 14 0 17 122 53	19 16 14 19 22 0 0 44 18	81 9 16 4 8 0 3 44 51	46 13 13 29 10 66 10 26 16	24 12 12 10 2 29 17 35 4 82	36 12 5 70 22 0 7 70 22
	SC.	ARLET	FEVI	ER CA	SE RA	TES		Sal		
98 cities	48	42	49	50	57	61	57	71	1 66	71
New England Middle Atlantic East North Central West North Central Bouth Atlantic East South Central West South Central Mountain Pacific	87 37 56 27 51 87 54 26 43	60 24 47 58 72 60 63 35 28	106 30 64 36 55 64 41 61 39	56 26 84 35 56 36 24 79 63	87 43 62 59 71 81 47 87 55	77 45 90 45 44 36 52 70 67	53 45 62 65 67 93 34 122 71	87 32 117 77 62 114 52 97 75	132 51 62 95 59 70 37 96 474	80 46 106 72 76 66 35 115 73
		SMALI	LPOX	CASE	RATES	3			- Sin	
98 cities	1	3	1	3	1	4	0	3	10	1
New England. Middle Atlantie. East North Central West North Central Gouth Atlantie. East South Central West South Central Mountain Pacific.	0 0 4 4 0 0 0 0	0 0 2 14 4 0 0 0 12	2 0 2 6 0 6 0	0 0 2 27 0 0 0 0 0 8	0 0 1 0 0 0 0 0 0	0 0 9 21 0 0 0 0	0 0 0 0 0 0 0	0 0 2 14 0 0 0 3 0 16	0 0 0 2 0 0 0 0 0	0 0 1 0 2 0 3 0

The figures given in this table are rates per 100,000 population, annual basis, and not the number of ases reported. Populations used are estimated as of July 1, 1931 and 1930, respectively.
 Waterloo, Iowa, and Spokane, Wash., not included.
 Waterloo, Iowa, not included.
 Spokane, Wash., not included.

Summary of weekly reports from cities, August 30 to October 3, 1931.—Annual rates per 100,000 population compared with rates for the corresponding period of 1930—Continued

TYPHOID FEVER CASE RATES

					Week e	ended-				
	Sept. 5, 1931	Sept. 6, 1930	Sept. 12, 1931	Sept. 13, 1930	Sept. 19, 1931	Sept. 20, 1930	Sept. 26, 1931	Sept. 27, 1930	Oet. 3, 1931	Oet. 4, 1930
98 cities	20	21	23	26	42	22	21	17	121	20
New England Middle Atlantic East North Central West North Central South Atlantic East South Central West South Central Mountain Pacific	7 13 16 6 49 41 74 44 10	12 20 12 14 58 48 45 9 8	7 13 10 13 79 35 91 35 27	22 24 17 21 70 48 52 62 4	22 16 91 38 26 47 44 26 35	12 15 11 29 68 48 63 0	5 16 15 36 43 47 47 26 10	12 13 9 15 56 18 35 44 12	17 21 9 14 65 52 24 26 14	12 14 9 14 42 60 52 115
	n	NFLUI	ENZA I	DEATI	RAT	ES				
91 cities	2	3	4	3	3	3	2	2	3	2
New England Middle Atlantic East North Central West North Central South Atlantic East South Central West South Central Mountain Pacific	2 1 1 3 2 6 10 0 2	0 3 2 6 8 0 11 9	2 4 3 9 2 0 17 0 2	0 4 3 0 2 19 0 0	2 3 3 6 4 0 0	2 2 2 0 0 26 7 18 0	0 1 3 0 4 6 0 0	2 2 2 0 4 13 4 0 5	2 3 2 12 0 6 0 0	13 11 18 2
	1	PNEUM	MONIA	DEAT	H RAT	ES				67
91 cities	50	53	55	54	60	57	52	57	53	58
New England Middle Atlantie East North Central West North Central. South Atlantie. East South Central West South Central Mountain Pacific.	24 62 33 62 61 38 83 96	56 65 36 51 68 91 50 53 27	58 65 36 44 63 82 73 70 46	68 63 43 45 58 26 57 123 25	50 66 45 44 57 57 57 93 78 84	56 65 42 75 56 71 46 115 40	67 55 38 44 51 32 52 70 86	39 72 47 36 56 65 71 53 40	58 60 35 59 61 63 66 61 58	44 59 53 69 52 104 71 183

Waterloo, Iowa, and Spokane, Wash., not included.
 Waterloo, Iowa, not included.
 Spokane, Wash., not included.

FOREIGN AND INSULAR

CANADA

Quebec Province—Communicable diseases—Week ended September 26, 1931.—The Bureau of Health of the Province of Quebec, Canada, reports cases of certain communicable diseases for the week ended September 26, 1931, as follows:

Disease	Cases	Disease	Cases
Chicken pox Diphtheria Erysipelas German measles Measles Mumps	36 47 2 3 16 11	Poliomyelitis. Scarlet fever. Tuberculosis. Typhoid fever Whooping cough.	105 47 39 23 24

CHINA

Shansi Province—Vital statistics—Year 1923.—According to the Nankai Weekly Statistical Service for July 13, 1931, published by the Institute of Economics of Nankai University at Tientsin, deaths from certain diseases occurred in the Province of Shansi during 1923 as shown in the table below. Evidently 1923 is the latest year for which such statistics for the Province have been published. The population in 1923 was given as 11,799,109.

Disease	Number of deaths	Death rate per 100,000 population	Disease	Number of deaths	Death rate per 100,000 population	
Cholera	2, 732 6, 647 7, 691 834	23. 2 65. 3 64. 4 7. 1	Measles Smallpox Tuberculosis	21, 625 8, 203 15, 108	183. 3 69. 5 128. 1	

The following table shows the number of births and deaths, the birth and death rates per 1,000 population, and the rate of natural increase in Shansi Province for the years 1912 to 1923:

	Bir	rths	De	og tie	
Year	Number	Rate per 1,000 pop- ulation	Number	Rate per 1,000 pop- ulation	Natural increase rate
1012 1913 1914 1915 1915 1916 1917 1918 1919 1920 1921	343, 015 327, 676 348, 648 448, 173 639, 988 703, 213 566, 153 145, 902 153, 035 150, 410 176, 634 180, 369	34. 0 32. 0 33. 4 43. 3 60. 8 62. 5 55. 7 12. 3 13. 4 12. 9 15. 1	218, 333 193, 791 142, 573 246, 534 421, 876 245, 503 242, 813 167, 374 132, 090 134, 977 160, 908 136, 709	21. 7 18. 9 13. 6 23. 8 40. 1 21. 7 23. 9 14. 1 11. 5 11. 6	12.3 13.1 19.8 20.7 40.8 31.8 -1.8 1.9

CUBA

Provinces—Communicable diseases—Four weeks ended August 29, 1931.—During the four weeks ended August 29, 1931, cases of certain communicable diseases were reported in the Provinces of Cuba, as follows:

Disease	Pinar del Rio	Habana	Matan-	Santa Clara	Cama- guey	Oriente	Total
Chicken pox. Diphtheria. Malaria Measles Paratyphold fover.		2 5 6 52	1 4	3 2 14	1	47 8 34	5. 16 46 70
Paratyphoid fever	3	23	10	47	9	29	12

DENMARK

Communicable diseases—August, 1931.—During the month of August, 1931, cases of certain communicable diseases were reported in Denmark as follows:

Disease	Cases	Disease	Cases
Anthrax Cerebrospinal meningitis Chicken pox Diphtheria and croup Erysipelas German measles Gonorrhea Influenza Lethargic encephalitis Measles	2 5 13 217 232 2 966 2,952 3 799	Mumps. Paratyphoid fever. Poliomyelitis. Scables	86 33 216 106 106 1,496

PANAMA CANAL ZONE

Communicable diseases—August, 1931.—During the month of August, 1931, certain communicable diseases, including imported cases, were reported in the Panama Canal Zone and terminal cities as follows:

Disease	Cases	Deaths	Disease	Cases	Deaths
Chicken pox Diphtheria Dysentery, amebic Dysentery, bacillary Leprosy Malaria	3 5 4	1	Measles. Mumps Pneumonia. Tuberculosis. Typhoid fever. Whooping cough.	44 1 10	a a

TASMANIA

Vital statistics—1930.—According to statistics published by the Commonwealth Bureau of Census and Statistics, at Hobart, Tasmania, births occurring during 1930 numbered 4,785 and deaths

1,948. There were 242 deaths of infants under 1 year of age, a rate of 50.6 per 1,000 births. The birth and death rates per 1,000 population in the urban and rural sections of Tasmania during the years 1920-1929, 1929, and 1930 are given in the accompanying table. The population of Tasmania in 1928 was approximately 215,000.

	1930	1929	1920-1929
Births per 1,000 population: Urban districts Rural districts Total	19. 3	19. 6	22. 7
	24. 4	24. 6	26. 2
	22. 2	22. 4	24. 8
Deaths per 1,000 population: Urban districts Rural districts Total	10.8	11. 4	11. 5
	7.7	9. 3	8. 5
	9.0	10. 2	9. 8

Cases of certain communicable diseases occurred in Tasmania during 1930, as compared with 1928 and 1929, as follows:

Disease	1930	1929	1928
Diphtheria Puerperal fever Scarlet fever Syphilis Tuberculosis Typhold fever	572	488	90
	27	25	2
	486	314	18
	26	34	2
	203	177	20
	27	49	8

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From medical officers of the Public Health Service, American consuls, International Office of Public Hygiene. Pan American Sanitary Bureau, health section of the League of Nations, and other sources. The reports contained in the following tables must not be considered as complete or final as regards either the list of countries included or the figures for which reports are given.

CHOLERA

[C indicates cases; D, deaths; P, present]

				-						Week ended-	-pape						
Place	Apr. 5- May 2, 1981	May 3-	May 31- June 27, 1931		July, 1981	1981			Упу	August, 1981	-		Ser.	September, 1931	r, 1931		October, 1931
					=	2	8	-		118	S	8	100	21	19	98	60
Ceylon: ColomboD						-											
Canton		64.	-		-		-	-	****	-	-	-	-	1	-	-	1
			22	1	1				-	•	-	-	8	•	98		
India.	120	13, 601		10°	2,848		-	4,029		4 11 2		\$	*	· · · · · ·	4		
-	176 21	265 149	108	Ea	22	లన్ వే	388	-433	02-	-8	250	024-	10000	0 M 00	- -		
Madrus		1.87	07	64	64			-		-61	00		-	-	-		
		i ca-			111-	09-	-		-	-		-					
Viragapatam India (French): O Mandermagor D Pandlebarro	80%	. :	4- 000	-			8		***	- 0101-						1 11	

CHOLERA, PLAGUE, SMALLPOX, TYPHUS FEVER, AND YELLOW FEVER-Continued

CHOLERA-Continued

[O indicates cases; D, deaths; P, present]

									4	Week ended-	-pop						
Place	Apr. 5- May 2, 1931		May 3- May 31- 30, June 1931 27, 1931		July, 1931	1881			Augr	August, 1931			Septe	September, 1931	1881	0	October, 1931
				-	п	18	8	-		15	23	83	12	2 19	8	60	10
India (Portuguese)	00		1		es-												
Indo-China (see also table below): Cochin-China—Rachgla			-		-	4-		А	-		-						
Saign and Cholon	*88 000	25	-24	00 PP	100	+0004			-					111-			
	9090										64-	-225°	1881				
	ADAC							100	G190	140	272 137 5	11	1.1	2922	\$238		2822
	POOD										9	01	60	11			00 00 0
ince	000000											8	288	1 2 2 2 2	128	182287	199
Suqelsbuyukh	ОДОДО	8										6161	19	111	111	- 20	•

Philippine Islands: 1 Provinces-

88	DODO			vessel: S. S. Arankola, at Rangoon from Calcutta. C S. S. Oity of Eastborne, at Calcutta from Co-		S.S. Cathay, at Kobe, Japan, from Shanghai,	S. S. Kasagi Maru, at Moji from Shanghal. C				e above):	
15		14 14 5	100	-					E.		00	100
44	នត	- +0		1						1961	125	883
	នន	1	1						March,	1831	188	322
	000								April,	1831	113	202
					0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0					1-10	88	848
		1	80 CH	1		N .			May, 1931	11-20	28	123
	-								-	21-31	9:	120
	-				6 6 6 6 9 9 9 9		0 0			1-10	23:	312
1	1	1	1			4.	*		June, 1931	11-20	8	282
3 10							-			21-30	120	
90							1 1	C4		1-10	22	8
5									July, 1931	11-20	8	30
\$13										21-31	56	849
12 16				-					Aug.	1931	30	*83

1 From May 3 to 25, 1831, 182 cases of cholera with 75 deaths were reported in Raisanjan and vicinity, Karman district, Persia.

1 Figures for cholera in the Philippine Islands are subject to correction.

1 Reports incomplete.

PLAGUE

IC indicates cases: D. deaths: P. present!

				-						-	Week ended	-pap						
Place	May 2,	2, 3-30,	May31- June 27,	27,	Ju	July, 1931	1	_		August, 1931	, 1931	1	-	Septe	September, 1931	1861	Octob	October, 1931
					11 +	18	32	2	8	15	22	81	2	13	8 19	98	00	10
Algeria: Algiera	0								64									
Bone Philippeville	000		11	1	9 9	1 1	11	11		11		1	11	11				
Argentina: San Juan Province. Belgian Congo.	000			1		A	I P			111	- 11							
British East Africa (see also table below): Tanganyika				1 11		1 1	9			+	1 00	+		-		-		
Ugsnda				1	95 132		986	1	1		100	11	11	11				
Ceylon: Colombo	100	2 40	200	980	- 1	-		277	2 4 4	8	8-1-	11	111		-		-	
Plague-infected rats.				•				-						-	•		1	1
Dutch East Indies:		ř	11	11	11	11	11	11	11	11	11	11	11	1 1				
Balavia and West Java.	000	¥1.	200	99	22	88	919	14	22	=======================================	88		99	11				
Java and Madura			19	102	59	22	52	99	38	23	74 47	11	90					
Kgypt: Alexandria	0	-	-	*	-	-	64	0.	*	100	-	1	64.	69	1	-		
Assiout	200	!	1001	* ='	1	1 1	1	•	-	79		1 1	- 1	-				
Beni-Suef		127	- 9	1		1 1	11	1 1	11		11	11			11	11		
Beheira	i	-	0	1 1	1 1	11	1 1		11	1 1	11	11	1 1	63		1 1	11	
Defrout	000	80	10	80		11	-		1	C4 .		11		11	11	11		
Gharbieh	900				11	11	11	-	1 1	1 1	11	11	1 1		11	11		
Oirps.		7	1	-		1 1	1 1		11		1 1	1 1		11	-	1	-	

Passes Passes Pague-infected rats Passes Passes Pague-infected rats Passes Pa										
d rats D D 6,142 752 88 11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		11	-	e4						
Pompenh C C C C C C C C C C C C C C C C C C C	_	8000	22 173 1171	83	215 137 1 1		00			
Phompenh	16	200	8 13	17	0	23 17	7	10	0	
Prompenh C	8	1,10				040	800			
	-04-04					01-1-	* -	N I I	N -	
gon a				-			5 - 1		01	
ovince.					100			-	1	

‡8

OA

PLAGUE-Continued

[C indicates cases; D, deaths; P, present]

												Week ended-	-pepu						
Place			Apr. 5- May 2, 1931	3-30, 1931	May31- June 27, 1931	1,1	July, 1931	1831			Augus	August, 1931			September, 1931	ber, 193		October,193	1881
						+	=	18	88	-	8	22 23	8	10	2	10	8		10
Tunisia: Tunis		00	91 8	91												-		1	
Union of South Africa: Cape Province		DA .					0 0 0 0 0 0 0 0 0 0 0 0												
Plague-infected rats. Orange Free State.		O	100	6464	1	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	1					-						III	111
Place	April, 1931	May, 1931	June, 1931	July, 1931	August, 1931	Septem- ber, 1931			Place	93		-	April, 1931	May, 1931	June, 1931	July, 1931	August, 1931	Bt, ter	Sep- tember, 1931
British East Africa (see also table above);							Peru					DA	00-	64	201		09		
a (see also table above). ar (see also table above)		-	2 6 55	\$	197		Sene	Senegal: Baol 1 Dakar 1	1			00000	64-1	4483	28		735.87	101 58 104 106	20 8 % % a
Antistrabe Province C Miarinarivo Province D Moramanos Province	\$7.00		22 -	23×				Louga 1Rufisque 1.	1 9			AOAOO	1	100414	4000		8-45	10-05	*****
Tananarive Province	48		20	10 10				Ilvaou	Tivaouane 1			AOA	7	119			1200-1	91	00

1 Reports incomplete.

SMALLPOY

SMALLPOX

									-	Week ended-	-pape						
Place	Apr. 5- May 2, 1931	May 3-30, 1931	May 31- June 27, 1931		July,	July, 1931			W	August, 1931	2		Bep	September, 1931	1831	-	Oct. 3.
					п	18	28	-	00	115	g	8	10	12	91	8	1931
Algeria: Algiers Constantine	000		œ 5	1		1							-				
gre (alastrim)	20 0	9		0	10	0	13	=		17		-	13				
British East Africa: Tanganyika	100	13	7	37	-8*		8		18			N			11	11	
British South Africa: Northern Rhodesia	00				. 2.												
	0 0		•		•			0 0 0	-			0 0			0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		
British Columbia. Manitoba	000		7				64	63			60		-	-	-		
Now innipeg Nowa Socia Ontario	000	17	25	7		9	12	-		2	64	-		- 69		111	
Kingston			L.				1						-	+	-		
South Ste. Marle. Toronto	000													-		0	
Quebec Saskatchewan Regina	200	\$ 44	25		13	10	10		10	9	10	œ	90	12	140		
Chile: Antologasta Chanaral	00	1		-							0 0 0 0 0 0 0 0						
China: Amoy																\pm	
Canton Foctow Hankow	000	on Pro	-A-4			64	-4		А		4-	-					
Hong Kong				1		•		•		1	•						: :

1 An epidemic of smallpox was reported on May 18 with 716 cases and 314 deaths since the middle of April, 1991, in Mendes Province, Bolivia,

SMALLPOX-Continued

[C indicates cases; D, deaths; P, present]

									-	Week ended-	-papu						
Place	Apr. 5- May 2, 1931	3-30, 1931	May31- June 27, 1931		Jul	July, 1931			'nγ	August, 1931	131		4	eptem	September, 1931		ő
				*	=	18	25	-	00	15	22	8	40	12	13	88	1931
China—Continued Manchuria— Harbin (see also table below).	DI	600					0 0		0 0 0 0 0					1 1	1 1	1	
Nanking Shanghal Foreigners only Including natives	A 128	Milit	1 12	6161		3					-	-			-		
Tientsin Chosen (see table below). Colombia:		64		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1					* : : : : : : : : : : : : : : : : : : :				1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1				
Santa Maria. Dutch East India: Batavia and West Java East Java and Medura.	00 00	21-												-			
Finland	000	80					24					98		1 1 1			
France (see table below). Great Britain: England and Wales. Bradford.			782	28	\$	88	98	8	88	19	2	33		10	45	45	
London London and Great Towns.	288°	183	122 228 1	53	• gg	34	32	e 2	202	17	10	13.3	38	11	32	35.01	
Sheffleld			-											-	-		

Tegucigalpa Tela. India.	0000 1,26 1-288	11,403	1, 704	1,492	1,376	1, 236	1, 255	828	203	645	88						1111
Bassein. Bombay. Calcutta.			2400	120	0+	*****	0000		74-	- -	1010-	0101		C4 C4C4			
Cochin Karachi Madras					64				•				1				1171
Negapatam Rangoon. Vizagapatam	00000A	82.001	******	8-8-		6	-	6-	n mm	-		m- mm	111	111		09	11111
India (French): Chandernagor Karikal			- 991	88	1111		1 8	11		900	000		000				11111
Pondicherry Province				**			381	==				- 1 1	•	-			
lon	0000						64	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	-						e4		
	0000	.0101				-					-					149	
Japan: Nagoya. Marico (see also table below): Jalisco (State)—Guadalajara. Mexico City and surrounding territory.	-22	431	-82	-27+		4-		00 17	64-	64		64	04		-	-	
Monterrey													-				

SMALLPOX-Continued

[C indicates cases; D, deaths; P, present]

									W	Week ended-	-per					
Place	Apr. 5- May 2, 1931	May 3-30, 1931	May31- June 27, 1931		July, 1931	1931			Augu	August, 1931	-		Ser	September, 1931	1, 1931	Oct. 3,
				*	=	10	8	-	•o	15	g	83	5 1	12 19	8	1931
Morocco (see table below). Poland Portugal: Lisbon Rumania (see table below).	266	679	ကတ္တေ့ မာ	15	1 22 1	18	-	14	23	x		10	12	8	-11	
Spain. Straits Settlements.		- 646	1			-1-		1 1 1	-		1 1 1	1 1 1				
	0 00	19 C	1					1 1		111		1 !		32		
Sudan (French) (see table below). Syla (see table below). Turkey (see table below). Union of Socialist Soviet Republics (see table below). Union of South Africa: Capp Province.	4	P.														
Natal Orange Pree State Transival Upper Volta	P 8	4 88	PP	24-	AA	44-				A	44	Ь	1 1 1 1	<u>a</u>		
On vessel: 8. S. Clan McTavish at Manila from Chittagong C 9. S. Talid (pilgrin ship) at Saukin from Jeddah C			1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0								1					

Place					Febru-	March,	April,		May, 1931	31		June, 1931	31		July, 1931	_	Y	August, 1931	1861
						1931	1031	1-10	11-20	21-31	1-10	11-20	21-30	1-10	11-20	21-81	1-10	11-20	21-31
Indo-Chine (see also table above) Ivory Coast Sudan (French).			0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	00000	35	82 4	142		11	4	8	16	-			14	11 28		
Place	Jan., 1931	Feb., 1931	Mar., 1931	Apr., 1931	May, 1931	June, 1931	July, 1931			Place	-		Jan., 1931	Feb., 1931	Mar., 1931	Apr., 1931	May, 1931	June, 1931	July, 1931
Chins: Harbin (see also table o			1		2	10		Turkey.	By			OA	83	37	1		6	1	
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[O indicates cases: D, deaths; P, present]

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TYPHUS PEVER-Continued

[C indicates cases: D, deaths; P, present]

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